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<p>(54) Title: MATERIALS AND METHODS FOR THE MODIFICATION OF ISOPRENOID CONTENT, COMPOSITION AND METABOLISM</p>		
<p>(57) Abstract</p> <p>Novel isolated polynucleotides associated with plant isoprenoid biosynthetic pathways are provided, together with genetic constructs comprising such sequences. Methods for the modulation of the content, structure and metabolism of polypeptides involved in an isoprenoid biosynthetic pathway in target organisms are also disclosed, the methods comprising incorporating one or more of the polynucleotides or genetic constructs of the present invention into the genome of a target organism. Modulation of the content, structure and metabolism of such polypeptides produces modifications in the content, structure and metabolism of isoprenoids in the target organism.</p>		

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## MATERIALS AND METHODS FOR THE MODIFICATION OF ISOPRENOID CONTENT, COMPOSITION AND METABOLISM

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### Technical Field of the Invention

This invention relates materials and methods for modifying the content, composition and metabolism of isoprenoids in plants and other organisms. More particularly, this invention relates to polypeptides involved in the synthesis of isoprenoid compounds, such as terpenoid and steroid compounds, polynucleotides encoding such polypeptides, expression of such polypeptides, and methods for modulating the composition and/or expression levels of such polypeptides, thereby modulating isoprenoid content, composition, and metabolism.

### 15 Background of the Invention

Isoprenoids form a large family of naturally occurring compounds, with over 20,000 distinct compounds having been described. The isoprenoids include vitamins A, D, E, and K, first recognized as fatty materials essential to the normal growth of animals, and numerous biological pigments. In plants, isoprenoid compounds, including terpenoid and steroid compounds, include hormones such as gibberellic acid and abscisic acid, pigments, electron carriers, membrane components (phytosterols), phytotoxins, antibiotics, flavors such as menthol, vitamins, macromolecular compounds such as rubber and guttapercha, and others.

Isoprene compounds, or prenyl lipids, are composed of one or more basic isoprene skeleton(s) ( $C_5$ ) formed by the decarboxylation of mevalonate-5-pyrophosphate. From the isopentenyl pyrophosphate ("active isoprene" or "IPP") and the isomeric dimethylallyl pyrophosphate, the geranyl pyrophosphate ( $C_{10}$ ) may be formed by "head-tail" condensation. By linkage of a further  $C_5$  unit, farnesyl pyrophosphate ( $C_{15}$ ) is formed. Further extension by "head-tail" or "tail-tail" condensation leads to  $C_{20}$ ,  $C_{30}$  and  $C_{40}$  compounds, as well as the higher molecular terpenoids. A schematic diagram of the basic biosynthetic pathways of isoprene compounds is shown in Fig. 1.

IPP is the branching point for a large variety of biologically significant molecules, including isoprenoids, carotenoids, and various sterols in different eukaryotic organisms

(mycosterols, phytosterols and zoosterols). In animals, cholesterol are precursors for several hormones and bile acids. Fungal ergosterol and mammalian cholesterol arise from IPP via squalene oxide and lanosterol, while higher plant sterols, like campesterol and sitosterol, are produced by cyclization of squalene oxide to cycloartenol and by further  
5 plant-specific enzymes.

Plant cells contain an intriguing diversity of a subclass of isoprenoids called terpenoids, most of which are cyclic with one or more rings. Terpenes in plants are divided into several classes, including sesquiterpenes, mono-, di-, triterpenes, etc. (Bohlmann *et al.*, *Proc. Natl. Acad. Sci. USA* 95:4126-4133, 1998). Terpenoids are formed by linking  
10 isoprene units ( $C_5H_8$ ) synthesized from acetate. Terpenoids include isoprene ( $C_5H_8$ ) compounds, including isopen-tenylpyrophosphate and active isoprene; monoterpene ( $C_{10}H_{16}$ ) compounds, including geraniol, and from which menthol, camphor, pinene and citronellal are derived; sesquiterpene ( $C_{15}H_{24}$ ) compounds, including farnesol, from which zingiberene, ubiquinone, plastoquinone, abscisic acid and rishitine are derived; diterpene  
15 ( $C_{20}H_{32}$ ) compounds, such as geranylgeraniol, from which phytol, kaurene, gibberellin acid and fusicoccin are derived; triterpene ( $C_{30}H_{48}$ ) compounds, including squalene, from which steroids and saponins are derived; tetraterpene ( $C_{40}H_{64}$ ) compounds, including phytoene and carotenes; and polyterpene ( $C_5H_8$ )<sub>n</sub> compounds, including rubber and guttapercha.

20 Synthase enzymes producing terpenes are thought to be of common evolutionary origin, lacking close similarity to other enzymes (except prenyltransferases). Most synthase enzymes have the ability to produce a variety of end-products from a single substrate. This may explain, in part, the enormous diversity of terpenoid compounds found in plants (Mitchell-Olds *et al.*, *Trends in Plant Science* 3(9):362-365, 1998). Complex  
25 terpene mixtures are thought to be important plant defensive compounds, their diversity and synergistic action delaying development of resistance in herbivores and pathogens (Langenheim J, *J. Chem. Ecol.* 20:1223-1280, 1994).

Plant terpenoids also have many known medicinal effects, and some plant isoprenoid compounds are administered as drugs. Taxol, which has proven to be  
30 efficacious in treating cancer, for example, is derived from terpenoid compounds. Dietary isoprenoids have been suggested to suppress mevalonate pathway, thereby affecting cancer and cardiovascular disease (Elson CE, *J. Nutr.* 125(6 Suppl):1666S-1672S, 1995). Farnesol, the last precursor common to all branches of the mevalonate pathway, has been

demonstrated to inhibit calcium channels in muscle cells (Roulette J-B, *J. Biol. Chem.* 51:32240-32246, 1997).

Ubiquinone and plastoquinone, which are also isoprenoid derivatives, function as electron carriers in the production of ATP in mitochondria and chloroplasts. In most mammalian tissues, ubiquinone (also called coenzyme Q) has ten isoprene units. Plastoquinone is the plant equivalent of ubiquinone. In their role as electron carriers, both ubiquinone and plastoquinone can accept either one or two electrons and either one or two protons to be reduced.

A remarkable role for isoprenyl intermediates has recently been discovered in studies of a protein that is implicated in human cancers and is known to associate with membranes through a covalently bound isoprenyl lipid. This protein, the RAS PROTEIN, is the product of the gene, a mutant version of a normal protein and a number of related GTP-binding proteins. The normal protein and the number of related GTP-binding proteins are known to act in signal transductions triggered by neurotransmitters, hormones, growth factors and other extracellular signals.

Quantitative and qualitative modifications in plant terpenoid content are known to be induced by external factors such as herbivore attack and wounding (Bohlmann J *et al.*, *Proc. Natl. Acad. Sci. USA* 95:6756-6761, 1998). Synthesis of cell terpenoids can also be induced by infection with pathogens. Even agricultural pest insects can be repelled by pine oil terpene compounds: monoterpenes carene, limonene and cymene deter onion flies (Ntiamoah Ya, *Entom. Exp. et Appl.* 79:219-226, 1996).

While the chemical diversity of isoprenoids is well known, and many of the metabolic pathways have been tentatively identified, few of the genes encoding enzymes responsible for the synthesis of isoprenoid compounds have been identified. The present invention is therefore directed to providing novel polynucleotides encoding polypeptides involved in the biosynthesis of isoprenoids, and providing methods for modifying the expression and composition of such polypeptides, thereby modulating isoprenoid content, composition, and metabolism.

Sequencing of the genomes, or portions of the genomes, of numerous biological materials, including humans, animals, microorganisms and various plant varieties, has been and is being carried out on a large scale. Polynucleotides identified using sequencing techniques may be partial or full-length genes, and may contain open reading frames, or portions of open reading frames, that encode polypeptides. Putative polypeptides may be

determined based on polynucleotide sequences. The sequencing data relating to polynucleotides thus represents valuable and useful information.

Polynucleotides may be analyzed for novelty by comparing identified sequences to sequences published in various public domain databases, such as EMBL. Newly  
5 identified polynucleotides and putative polypeptides may also be compared to polynucleotides and polypeptides contained in databases to ascertain homology to known polynucleotides and polypeptides. In this way, the degree of similarity or identity or homology of polynucleotides and polypeptides having an unknown function may be determined relative to polynucleotides and polypeptides having known functions.

10 U.S. Patent 5,589,619 discloses materials and methods for increasing squalene and sterol accumulation in higher plants by modifying the copy number of a gene encoding a polypeptide having HMG-CoA reductase activity. Genetic materials, including polynucleotides, polypeptides, DNA molecules, and the like, relating to HMG-CoA reductase activity are disclosed, as well as methods for transforming plant cells and  
15 producing transgenic plants.

U.S. Patent 5,689,047 discloses stilbene synthase genes derived from grapevines, as well as the use of those genes in vectors and transformed microorganisms, as well as transformed plant cells and plants.

U.S. Patent 5,753,507 discloses plant polynucleotides encoding geraniol/nerol 10 –  
20 hydroxylase ( $G_{10}H$ ), as well as methods for using complete and partial polynucleotides as probes, and methods for expressing  $G_{10}H$  and enhancing levels of terpenoid indole alkaloid and iivoid insect pheromone produced by a plant.

The following U.S. Patents disclose isoprenoid compounds or related compounds, or methods for using such compounds: U.S. Patent 5,429,939; U.S. Patent 5,444,166; U.S.  
25 Patent 5,460,949; U.S. Patent 5,470,832; U.S. Patent 5,474,925; U.S. Patent 5,495,070; U.S. Patent 5,521,078; U.S. Patent 5,545,816; U.S. Patent 5,547,856; U.S. Patent 5,569,832; U.S. Patent 5,580,963; U.S. Patent 5,597,718; U.S. Patent 5,670,349; U.S. Patent 5,674,485; U.S. Patent 5,684,238; U.S. Patent 5,689,056; U.S. Patent 5,691,147; U.S. Patent 5,693,476; and U.S. Patent 5,443,978. The U.S. Patents cited above are  
30 incorporated by reference herein in their entireties.

### Summary of the Invention

Briefly, the present invention provides isolated polynucleotides encoding polypeptides involved in the production and modification of isoprenoids. Genetic constructs comprising such sequences and methods for the use of such genetic constructs  
5 are also provided, together with transgenic cells and plants incorporating such genetic constructs and exhibiting modified isoprenoid content, composition, and metabolism.

In a first aspect, the present invention provides isolated polynucleotide sequences identified in the attached Sequence Listing as SEQ ID NOS: 1-53 and 78-164, variants of those sequences, extended sequences comprising the sequences set out in SEQ ID NOS:  
10 1-53, 78-164 and their variants, probes and primers corresponding to the sequences set out in SEQ ID NOS: 1-53, 78-164 and their variants, polynucleotides comprising at least a specified number of contiguous residues of any of the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164 (x-mers), and extended sequences comprising portions of the sequences set out in SEQ ID NOS: 1-53 and 78-164, all of which are referred to herein,  
15 collectively, as "polynucleotides of the present invention."

The present invention also provides isolated polypeptide sequences identified in the attached Sequence Listing as SEQ ID NOS: 165-304, polypeptide variants of those sequences, polypeptides comprising the isolated polypeptide sequences and variants of those sequences, polypeptides comprising at least a specified number of contiguous  
20 residues of any of the polypeptides identified as SEQ ID NOS: 165-304; and polypeptides comprising portions of the sequences set out in SEQ ID NOS: 165-304.

The polynucleotide sequences identified as SEQ ID NOS: 1-53 and 78-164 were derived from plant sources, namely from *Eucalyptus grandis* and *Pinus radiata*. The polynucleotides of the present invention are primarily "partial" sequences, in that they do  
25 not represent a full length gene encoding a full length polypeptide. Such partial sequences may be extended by analyzing and sequencing various DNA libraries using primers and/or probes and well known hybridization and/or PCR techniques. The partial sequences identified as SEQ ID NOS: 1-53 and 78-164 may thus be extended until an open reading frame encoding a polypeptide, a full length polynucleotide and/or gene capable of  
30 expressing a polypeptide, or another useful portion of the genome is identified. Such extended sequences, including full length polynucleotides and genes, are described as "corresponding to" a sequence identified as one of the sequences of SEQ ID NOS: 1-53 and 78-164 or a variant thereof, or a portion of one of the sequences of SEQ ID NOS: 1-53

and 78-164 or a variant thereof, when the extended polynucleotide comprises an identified sequence or its variant, or an identified contiguous portion (x-mer) of one of the sequences of SEQ ID NOS: 1-53 and 78-164 or a variant thereof. Similarly, RNA sequences, reverse sequences, complementary sequences, anti-sense sequences, and the like, corresponding to  
5 the polynucleotides of the present invention, may be routinely ascertained and obtained using the cDNA sequences identified as SEQ ID NOS: 1-53 and 78-164.

The polynucleotides identified as SEQ ID NOS: 1-53 and 78-164 may contain open reading frames ("ORFs") or partial open reading frames encoding polypeptides. Additionally, open reading frames encoding polypeptides may be identified in extended or  
10 full length sequences corresponding to the sequences set out as SEQ ID NOS: 1-53 and 78-164. Open reading frames may be identified using techniques that are well known in the art. These techniques include, for example, analysis for the location of known start and stop codons, most likely reading frame identification based on codon frequencies, etc. Suitable tools and software for ORF analysis are available, for example, on the Internet at  
15 <http://www.ncbi.nlm.nih.gov/gorf/gorf.html>. Open reading frames and portions of open reading frames may be identified in the polynucleotides of the present invention. Once a partial open reading frame is identified, the polynucleotide may be extended in the area of the partial open reading frame using techniques that are well known in the art until the polynucleotide for the full open reading frame is identified. Thus, polynucleotides and  
20 open reading frames encoding polypeptides may be identified using the polynucleotides of the present invention.

Once open reading frames are identified in the polynucleotides of the present invention, the open reading frames may be isolated and/or synthesized. Expressible DNA constructs comprising the open reading frames and suitable promoters, initiators,  
25 terminators, etc., which are well known in the art, may then be constructed. Such DNA constructs may be introduced into a host cell to express the polypeptide encoded by the open reading frame. Suitable host cells may include various prokaryotic and eukaryotic cells, including plant cells.

Polypeptides encoded by the polynucleotides of the present invention may be  
30 expressed and used in various assays to determine their biological activity. Such polypeptides may be used to raise antibodies, to isolate corresponding interacting proteins or other compounds, and to quantitatively determine levels of interacting proteins or other compounds.



The present invention also contemplates methods for modulating the polynucleotide and/or polypeptide content and composition of an organism, such methods involving, according to one embodiment, stably incorporating into the genome of the organism a genetic construct comprising one or more polynucleotides of the present invention. In one embodiment, the target organism is a plant, preferably a woody plant, more preferably a woody plant of the *Pinus* or *Eucalyptus* species, and most preferably *Eucalyptus grandis* or *Pinus radiata*. In a related aspect, a method for producing an organism having an altered genotype or phenotype is provided, the method comprising transforming a host cell with a genetic construct of the present invention to provide a transgenic cell, and cultivating the transgenic cell under conditions conducive to growth and regeneration. Organisms having an altered genotype or phenotype as a result of modulation of the level or content of a polynucleotide or polypeptide of the present invention compared to a wild-type organism, as well as components (seeds, etc.) of such organisms and progeny of such organisms, are contemplated by and encompassed within the present invention.

The isolated polynucleotides of the present invention have utility in genome mapping, in physical mapping, and in positional cloning of genes. Additionally, the polynucleotide sequences identified as SEQ ID NOS: 1-53, 78-164, and their variants, may be used to design oligonucleotide probes and primers. Oligonucleotide probes and primers have sequences that are substantially complementary to the polynucleotide of interest over a certain portion of the polynucleotide. Oligonucleotide probes designed using the polynucleotides of the present invention may be used to detect the presence and examine the expression patterns of genes in any organism having sufficiently similar DNA and RNA sequences in their cells using techniques that are well known in the art, such as slot blot DNA hybridization techniques. Oligonucleotide primers designed using the polynucleotides of the present invention may be used for PCR amplifications. Oligonucleotide probes and primers designed using the polynucleotides of the present invention may also be used in connection with various microarray technologies, including the microarray technology used by Synteni (Palo Alto, CA).

The polynucleotides of the present invention may also be used to tag or identify an organism or reproductive material therefrom. Such tagging may be accomplished, for example, by stably introducing a non-disruptive non-functional heterologous

polynucleotide identifier into an organism, the polynucleotide comprising one of the polynucleotides of the present invention.

The polynucleotides of the present invention encode polypeptides that have activity in an isoprenoid biosynthetic pathway. The isoprenoid metabolism-related polynucleotides were isolated from pine and eucalyptus, and putatively identified by DNA and protein similarity searches. Various isoprenoid compounds are well characterized and have useful properties. Methods of the present invention relating to modulating the polynucleotide and/or polypeptide content and composition of an organism and, thereby, modulating the isoprenoid content, composition and metabolism of an organism, are applicable to a wide range of activities. The novel materials and methods of the present invention have a multitude of potential uses: in forestry and agriculture for manipulation of isoprenoid metabolism; in medicine for therapeutic effects, including direct application in diseased organisms or indirect application by transgenic organisms; in fermentation and chemical processing industries involving isoprenoids; and in numerous other applications, some of which are described in the references cited above. In plant applications, manipulating isoprenoid pathways or isoprenoid composition may, for example, affect plant development, pest resistance, and the value of extractives (pinene, myrcene, etc.). In foodstuffs, various isoprenoids affect the nutritional quality and pharmacological properties of the ingested material, e.g. cholesterol or phytosterol composition of animal-derived and plant-derived foods for human or animal consumption. Additionally, isoprenoid pathways control the production of vitamins A, E, and K; plant pigments such as carotene and the phytol chain of chlorophyll; natural rubber; many essential oils, such as the fragrant principles of lemon oil, eucalyptus, and musk; insect juvenile hormone, which controls metamorphosis; dolichols, which serve as lipid-soluble carriers in complex polysaccharide synthesis; and ubiquinone and plastoquinone, electron carriers in mitochondria and chloroplasts. The ubiquitous and varied roles of isoprenoids thus make these compounds and the polynucleotides encoding them attractive targets for biotechnical applications in a variety of fields.

Briefly, the present invention provides isolated polynucleotides encoding polypeptides involved in the synthesis of isoprenoids. The polynucleotides and polypeptides of the present invention have demonstrated similarity to polypeptides that are known to be involved in the synthesis of isoprenoids as shown below in Table 1.

TABLE 1

POLYNUCLEOTIDE SEQ ID NO	POLYPEPTIDE SEQ. ID	POLYPEPTIDE IDENTITY
1	252	Acetylcholinesterase Precursor
2	253	Deoxyxylulosephosphate Synthase (DXPS)
3, 4, 44	254,255,295	Geranyltranstransferase
5, 6	256,266	Farnesyltranstransferase
7, 154	258 241	Squalene Synthetase
8-10, 155-157	259-261 242-244	Squalene Monooxygenase
11	262	Geranylgeranyl-Diphosphate Geranylgeranyltransferase
12	263	Trichodiene Synthase
13, 25, 84-88, 95 115-118	264,276 171-175, 182, 202-205	Pinene Synthase
14, 89, 90	265 176, 177	Abietadine Synthase
15, 91-94, 96-98, 131-135	266 178-181, 183-185, 218-222	Hydroxymethylglutaryl-Coa Reductase (NADPH)
16, 17, 18, 99-102	267,268,269, 186-189	Myrcene Synthase
19, 20, 103, 107, 108	270,271 190, 194, 195	Limonene Synthase
21-23, 109-111	272-274 196-198	Cadinene Synthase
24, 114	275 201	Bisabolene Synthase
26, 27	277,278	Pinene/Myrcene/Limonene Synthase
28, 119-122	279 206-209	Cycloartenol Synthase
29, 124-126	280 211-213	Obtusifoliol Demethylase
30	281	Lupeol Synthase
31, 158, 159	282 245, 246	Udp-Glucose:Sterol Glucosyltransferase
32	283	Hydroxymethylglutaryl-CoA Reductase (NADPH)
33, 34, 160-162	284,285 247-249	Sterolmethyltransferase
35, 136	286 223	Lecithin:Cholesterol Acyl Transferase
36, 137	287 224	Sterol Delta-7 Reductase
37, 38, 138-140	288,289 225-227	Methyl Sterol Oxidase
39	290	Deoxyxylulosephosphate Synthase (DXPS)
40	291	Phosphomevalonate Kinase
41, 50, 141, 142, 146	292,301 228, 229, 233	Diphosphomevalonate Decarboxylase
42, 43, 143	293,294 230	Isopentenyl-Diphosphate Delta- Isomerase

POLYNUCLEOTIDE SEQ ID NO	POLYPEPTIDE SEQ. ID	POLYPEPTIDE IDENTITY
45	296	Estradiol Dehydrogenase
46-49, 144, 145	297-300 231, 232	Furostanol Glucosidase
51, 52, 147-153	302,303 234-240	Oxysterol-Binding Protein
53	304	Sterol Carrier Protein
78, 79, 127-130	165, 166, 214-217	Sterol 14-demethylase
81	168	Sesquiterpene cyclase
82, 83	169, 170	Geranylgeranyl diphosphate
104-106, 164	191-193, 251	CXPS/transketolase
112, 113	199, 200	Sabinene synthase
123	210	Beta-amyrin synthase
163	250	Sterol desaturase

In one embodiment, the isolated polynucleotides comprise a sequence selected from the group consisting of: (a) sequences recited in SEQ ID NOS: 1-53 and 78-164;  
 5 (b) complements of the sequences recited in SEQ ID NOS: 1-53 and 78-164; (c) reverse complements of the sequences recited in SEQ ID NOS: 1-53 and 78-164; (d) reverse sequences of the sequences recited in SEQ ID NOS: 1-53 and 78-164; and (e) sequences having either 40%, 60%, 75% or 90% identity, as defined herein, to a sequence of (a) – (d) or a specified region of a sequence of (a) – (d).

10 In a further aspect, isolated polypeptides encoded by the polynucleotides of the present invention are provided. In one embodiment, such polypeptides comprise an amino acid sequence encoded by polynucleotides of the present invention, including polynucleotides comprising a sequence set out in the group consisting of SEQ ID NOS: 1-53 and 78-164, as well as polypeptides comprising an amino acid sequence recited in SEQ  
 15 ID NOS: 165- 304.

In another aspect, the invention provides genetic constructs comprising a polynucleotide of the present invention, either alone, in combination with one or more additional polynucleotides of the present invention, or in combination with one or more known polynucleotides, together with transgenic cells comprising such constructs.

20 In a related aspect, the present invention provides genetic constructs comprising, in the 5'-3' direction, a gene promoter sequence; an open reading frame coding for at least a functional portion of an enzyme encoded by an inventive polynucleotide or a variant thereof; and a gene termination sequence. The open reading frame may be oriented in either a sense or antisense direction. Genetic constructs comprising a non-coding region

of a gene coding for an enzyme encoded by the above polynucleotide or a nucleotide sequence complementary to a non-coding region, together with a gene promoter sequence and a gene termination sequence, are also provided. Genetic constructs comprising, in the 5' – 3' direction, a promoter sequence; a polynucleotide sequence comprising at least one  
5 of the following: (1) a polynucleotide comprising a polynucleotide of the present invention; or (2) a polynucleotide comprising a polynucleotide of the present invention and including a non-coding region of a gene coding for a polypeptide having activity in an isoprenoid biosynthetic pathway, are also contemplated. The genetic construct may further include a marker for the identification of transformed cells.

10 In a further aspect, transgenic host cells, such as transgenic plant cells, comprising the genetic constructs of the present invention are provided, together with plants comprising such transgenic cells, and fruits, seeds, and progeny of such plants. Other useful host cells include bacterial cells, insect cells, yeast cells and mammalian cells.

In yet another aspect, methods for modulating the isoprenoid content, composition,  
15 and metabolism of an organism are provided, such methods including stably incorporating into the genome of the organism a genetic construct of the present invention. In a preferred embodiment, the target organism is a plant and the plant is a woody plant, preferably selected from the group consisting of eucalyptus, pine, acacia, poplar, sweetgum, teak and mahogany species, more preferably from the group consisting of pine  
20 and eucalyptus species, and most preferably from the group consisting of *Eucalyptus grandis* and *Pinus radiata*. In a related aspect, a method for producing an organism having modified isoprenoid content is provided, the method comprising transforming a host cell with a genetic construct of the present invention to provide a transgenic cell and cultivating the transgenic cell under conditions conducive to growth and regeneration.

25 In yet a further aspect, the present invention provides methods for modifying the activity of a polypeptide in a target organism such as a plant, comprising stably incorporating into the genome of the organism a genetic construct of the present invention. In a preferred embodiment, the target organism is a plant, and the plant is a woody plant, preferably selected from the group consisting of eucalyptus, pine, acacia, poplar,  
30 sweetgum, teak and mahogany species, more preferably from the group consisting of pine and eucalyptus species, and most preferably from the group consisting of *Eucalyptus grandis* and *Pinus radiata*.

In yet a further aspect, the present invention provides methods for modulating one or more of the content, the composition and the metabolism of an isoprenoid compound in an organism by administering an isolated polypeptide of the present invention to the organism. In applications in which the organism is a plant, administration of the  
5 polypeptide may be topical, such as by spraying or similar topical application. In applications in which the organism is mammalian, administration of the polypeptide may be systemic, such as by injection, intradermal delivery, oral delivery, delivery via nasal passageways or airways, or the like.

The above-mentioned and additional features of the present invention and the  
10 manner of obtaining them will become apparent, and the invention will be best understood by reference to the following more detailed description.

### Description of Drawings

Fig. 1 shows a schematic diagram illustrating basic biosynthetic pathways of  
15 isoprene compounds.

Fig. 2 illustrates genomic DNA samples from tobacco plants created in a tagging experiment using a unique sequence identifier from *Pinus* (left panel) and a unique sequence identifier from *Eucalyptus* (right panel). In both panels, Lanes A and B contain DNA samples from empty-vector transformed control plants and Lanes C-E contain DNA  
20 samples from plants transformed with a unique sequence identifier.

Fig. 3 illustrates detection of a *Pinus* unique sequence identifier in transformed tobacco plants. Lanes A and B show the hybridization of a probe from SEQ ID NO: 76 to the genomic DNA of tobacco plants which lack the *Pinus* unique sequence identifier (empty-vector transformed control plants). Lanes C-E show the hybridization of the probe  
25 to the genomic DNA of tobacco plants containing one to three copies of the *Pinus* unique sequence identifier.

Fig. 4 illustrates detection of a *Eucalyptus* unique sequence identifier in transformed tobacco plants. Lanes A and B show the hybridization of a probe from SEQ ID NO: 77 to the genomic DNA of tobacco plants which lack the *Eucalyptus* unique  
30 sequence identifier (empty-vector transformed control plants). Lanes C-E show the hybridization of the probe to the genomic DNA of tobacco plants containing one to two copies of the *Eucalyptus* unique sequence identifier.

### Detailed Description

As described above, isoprenoids are important components in a variety of eukaryotic functions. Modification of isoprenoid content, composition, and metabolism in the earlier parts of the pathway, especially the steps up to the formation of isopentenyl-diphosphate (IPP), geranyl-diphosphate (GPP), farnesyl-diphosphate (FPP) and squalene, may have a profound influence on the synthesis of the isoprenoid compounds deriving from these two precursors. Blocking one or more of the downstream steps branching from isopentenyl-diphosphate and squalene may also have a substantial effect on the pool of isopentenyl-diphosphate and squalene available for synthesis of terpenes or steroids. Hence, modifications in the synthesis, content, composition, and metabolism of any single enzyme in the isoprenoid biosynthetic pathway, and particularly in the early part of the pathway (IPP => GPP => FPP => squalene) of the isoprenoid synthesis, may affect the content, composition and metabolism of terpenoid and steroid compounds.

Using the methods and materials of the present invention, the isoprenoid content of a plant may be modified by incorporating sense or antisense copies of polynucleotides encoding polypeptides involved in the synthesis of isoprenoids into the genome of a target organism. In addition, the number of copies and combination of polynucleotides encoding for different enzymes in the biosynthetic pathway of isoprenoids may be manipulated to modify the relative amounts of isoprenoids synthesized, thereby producing biological materials having an altered composition and/or altered isoprenoid metabolism. Similarly, the alteration of isoprenoid composition, for direct application in a target organism, or for production of polypeptides for separate use, is advantageous for a variety of applications, as evidenced by the references cited above and incorporated herein by reference.

According to one embodiment, the present invention provides isolated polynucleotides encoding, or partially encoding, polypeptides having similarity to polypeptides known to be involved in isoprenoid synthesis and modification. The polynucleotides of the present invention were isolated from eucalyptus and pine species, but may alternatively be isolated from other plant sources and may be synthesized using conventional synthesis techniques. Specifically, isolated polynucleotides of the present invention comprise: the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164; complements of the sequences identified as SEQ ID NOS: 1-53 and 78-164; reverse sequences of the sequences identified as SEQ ID NOS: 1-53 and 78-164; reverse complements of the sequences identified as SEQ ID NOS: 1-53 and 78-164; at least a

specified number of contiguous residues (*x*-mers) of any of the above-mentioned polynucleotides; polynucleotides complementary to any of the above polynucleotides; anti-sense sequences corresponding to any of the above polynucleotides; and variants of any of the above polynucleotides, as that term is described in this specification.

5        The isolated polynucleotides recited in SEQ ID NOS: 1-53 and 78-164 encode, or partially encode, polypeptides demonstrating sequence similarity to polypeptides known to be involved in an isoprenoid biosynthetic pathway, as indicated in Table 1 above. More specifically, the isolated polynucleotides listed in the first column of Table 1 encode, or partially encode the polypeptides listed in alignment in the second column of Table 1,  
10    above. Predicted amino acid sequences corresponding to the polynucleotides set out in SEQ ID NOS: 1-53, 78-164, based on information available at the time of filing this application, are provided in SEQ ID NOS: 165-304, as indicated in Table 1.

      The term "polynucleotide(s)," as used herein, means a single or double-stranded polymer of deoxyribonucleotide or ribonucleotide bases and includes DNA and  
15    corresponding RNA molecules, including HnRNA and mRNA molecules, both sense and anti-sense strands, and comprehends cDNA, genomic DNA and recombinant DNA, as well as wholly or partially synthesized polynucleotides. An HnRNA molecule contains introns and corresponds to a DNA molecule in a generally one-to-one manner. An mRNA molecule corresponds to an HnRNA and DNA molecule from which the introns have been  
20    excised. A polynucleotide may consist of an entire gene, or any portion thereof. A gene is a polypeptide that codes for a functional polypeptide or RNA molecule. Operable anti-sense polynucleotides may comprise a fragment of the corresponding polynucleotide, and the definition of "polynucleotide" therefore includes all such operable anti-sense fragments. Anti-sense polynucleotides and techniques involving anti-sense  
25    polynucleotides are well known in the art and are described, for example, in Robinson-Benion *et al.*, *Methods in Enzymol.* 254(23):363-375, 1995; and Kawasaki *et al.*, *Artific. Organs* 20(8):836-848, 1996. Polynucleotides of the present invention also encompass polynucleotide sequences that differ from the disclosed sequences but which, as a result of the degeneracy of genetic code, encode a polypeptide which is the same as that encoded  
30    by a polynucleotide of the present invention.

      The definitions of the terms "complement," "reverse complement," and "reverse sequence," as used herein, are best illustrated by the following examples. For the



sequence 5' AGGACC 3', the complement, reverse complement, and reverse sequences are as follows:

	complement	3' TCCTGG 5'
	reverse complement	3' GGTCCT 5'
5	reverse sequence	5' CCAGGA 3'

Identification of genomic DNA and heterologous species DNAs can be accomplished by standard DNA/DNA hybridization techniques, under appropriately stringent conditions, using all or part of a cDNA sequence as a probe to screen an appropriate library. Alternatively, PCR techniques using oligonucleotide primers that are designed based on known genomic DNA, cDNA and protein sequences can be used to amplify and identify genomic and cDNA sequences. Synthetic DNAs corresponding to the identified sequences and variants may be produced by conventional synthesis methods. All of the polynucleotides described herein are isolated and purified, as those terms are commonly used in the art.

In another aspect, the present invention provides isolated polypeptides encoded, or partially encoded, by the above polynucleotides. As used herein, the term "polypeptide" encompasses amino acid chains of any length, including full length proteins, wherein the amino acid residues are linked by covalent peptide bonds. The term "polypeptide encoded by a polynucleotide" as used herein, includes polypeptides encoded by a polynucleotide which comprises an isolated polypeptide or variant provided herein. In one embodiment, polypeptides of the present invention comprise an amino acid sequence selected from the group consisting of sequences provided in SEQ ID NOS: 165-304, as well as variants of such sequences. According to another embodiment, polypeptides of the present invention comprise at least a specified number of contiguous residues (x-mers) of any of the sequences provided in SEQ ID NOS: 165-304.

Polypeptides of the present invention may be produced recombinantly by inserting a polynucleotide that encodes the polypeptide into an expression vector and expressing the polypeptide in an appropriate host. Any of a variety of expression vectors known to those of ordinary skill in the art may be employed. Expression may be achieved in any appropriate host cell that has been transformed or transfected with an expression vector containing a polypeptide encoding a recombinant polypeptide. Suitable host cells include prokaryotes, yeast and higher eukaryotic cells. Preferably, the host cells employed are *Escherichia coli*, insect, yeast or a mammalian cell line such as COS or CHO. The

polynucleotide(s) expressed in this manner may encode naturally occurring polypeptides, portions of naturally occurring polypeptides, or other variants thereof.

In a related aspect, polypeptides are provided that comprise at least a functional portion of a polypeptide having an amino acid sequence selected from the group consisting of sequences provided in SEQ ID NOS: 165-304, and variants thereof. As used herein, a  
5 "functional portion" of a polypeptide is that portion which contains the active site essential for affecting the function of the polypeptide, for example, the portion of the molecule that is capable of binding one or more reactants. The active site may be made up of separate portions present on one or more polypeptide chains and will generally exhibit high binding  
10 affinity.

Functional portions of a polypeptide may be identified by first preparing fragments of the polypeptide by either chemical or enzymatic digestion of the polypeptide, or by mutation analysis of the polynucleotide that encodes the polypeptide and subsequent expression of the resulting mutant polypeptides. The polypeptide fragments or mutant  
15 polypeptides are then tested to determine which portions retain biological activity, using, for example, the representative assays provided below.

A functional portion comprising an active site may be made up of separate portions present on one or more polypeptide chains and generally exhibits high substrate specificity. The term "polypeptide encoded by a polynucleotide" as used herein, includes  
20 polypeptides encoded by a polynucleotide comprising a partial isolated polynucleotide of the present invention.

Portions and other variants of the inventive polypeptides may also be generated by synthetic or recombinant means. Synthetic polypeptides having fewer than about 100 amino acids, and generally fewer than about 50 amino acids, may be generated using  
25 techniques that are well known to those of ordinary skill in the art. For example, such polypeptides may be synthesized using any of the commercially available solid-phase techniques, such as the Merrifield solid-phase synthesis method, where amino acids are sequentially added to a growing amino acid chain. See Merrifield, *J. Am. Chem. Soc.* 85:2149-2154, 1963. Equipment for automated synthesis of polypeptides is commercially  
30 available from suppliers such as Perkin Elmer/Applied Biosystems, Inc. (Foster City, CA), and may be operated according to the manufacturer's instructions. Variants of a native polypeptide may be prepared using standard mutagenesis techniques, such as oligonucleotide-directed site-specific mutagenesis (Kunkel T, *Proc. Natl. Acad. Sci. USA*

82: 488-492, 1985). Sections of DNA sequences may also be removed using standard techniques to permit preparation of truncated polypeptides.

In general, the polypeptides disclosed herein are prepared in an isolated, substantially pure form. Preferably, the polypeptides are at least about 80% pure; more preferably at least about 90% pure; and most preferably, at least about 99% pure. In certain preferred embodiments, described in detail below, the isolated polypeptides are incorporated into pharmaceutical compositions or vaccines for use in the treatment of skin disorders.

As used herein, the term "variant" comprehends polynucleotide or polypeptide sequences different from the specifically identified sequences, wherein one or more nucleotides or amino acid residues is deleted, substituted, or added. Variants may be naturally occurring allelic variants, or non-naturally occurring variants. Variant polynucleotide sequences preferably exhibit at least 40%; more preferably at least 60%; more preferably yet at least 75%; and most preferably at least 90% identity to a sequence of the present invention. Variant polypeptide sequences preferably exhibit at least 50%; more preferably at least 75%; more preferably yet at least 90%; and most preferably at least 95% identity to a sequence of the present invention. The percentage identity is determined by aligning the two sequences to be compared as described below, determining the number of identical residues in the aligned portion, dividing that number by the total number of residues in the inventive (queried) sequence, and multiplying the result by 100.

Polynucleotide and polypeptide sequences may be aligned, and percentage of identical residues in a specified region may be determined against another polynucleotide or polypeptide, using computer algorithms that are publicly available. Two exemplary algorithms for aligning and identifying the similarity of polynucleotide sequences are the BLASTN and FASTA algorithms. Polynucleotides may also be analyzed using the BLASTX algorithm, which compares the six-frame conceptual translation products of a nucleotide query sequence (both strands) against a protein sequence database. The percentage identity of polypeptide sequences may be examined using the BLASTP algorithm. The BLASTN, BLASTX and BLASTP programs are available on the NCBI anonymous FTP server (<ftp://ncbi.nlm.nih.gov>) under /blast/executables/. The BLASTN algorithm Version 2.0.4 [Feb-24-1998] and Version 2.0.6 [Sept-16-1998], set to the parameters described below, is preferred for use in the determination of polynucleotide variants according to the present invention. The BLASTP algorithm, set to the parameters

described below, is preferred for use in the determination of polypeptide variants according to the present invention. The use of the BLAST family of algorithms, including BLASTN, BLASTP, and BLASTX, is described at NCBI's website at URL <http://www.ncbi.nlm.nih.gov/BLAST/newblast.html> and in the publication of Altschul, *et al.*, *Nucleic Acids Res.* 25: 3389-3402, 1997.

The computer algorithm FASTA is available on the Internet at the ftp site <ftp://ftp.virginia.edu/pub/fasta/>. Version 2.0u4 [February 1996], set to the default parameters described in the documentation and distributed with the algorithm, may be also used in the determination of variants according to the present invention. The use of the FASTA algorithm is described in Pearson and Lipman, *Proc. Natl. Acad. Sci. USA* 85:2444-2448, 1988; and Pearson WR, *Methods in Enzymol.* 183: 63-98, 1990.

The following running parameters are preferred for determination of alignments and similarities using BLASTN that contribute to the E values and percentage identity for polynucleotide sequences: Unix running command: `blastall -p blastn -d embldb -e 10 -G0 -E0 -r 1 -v 30 -b 30 -i queryseq -o results`; the parameters are: -p Program Name [String]; -d Database [String]; -e Expectation value (E) [Real]; -G Cost to open a gap (zero invokes default behavior) [Integer]; -E Cost to extend a gap (zero invokes default behavior) [Integer]; -r Reward for a nucleotide match (BLASTN only) [Integer]; -v Number of one-line descriptions (V) [Integer]; -b Number of alignments to show (B) [Integer]; -i Query File [File In]; and -o BLAST report Output File [File Out] Optional.

The following running parameters are preferred for determination of alignments and similarities using BLASTP that contribute to the E values and percentage identity of polypeptide sequences: `blastall -p blastp -d swissprot -e 10 -G 0 -E 0 -v 30 -b 30 -i queryseq -o results`; the parameters are: -p Program Name [String]; -d Database [String]; -e Expectation value (E) [Real]; -G Cost to open a gap (zero invokes default behavior) [Integer]; -E Cost to extend a gap (zero invokes default behavior) [Integer]; -v Number of one-line descriptions (v) [Integer]; -b Number of alignments to show (b) [Integer]; -i Query File [File In]; -o BLAST report Output File [File Out] Optional. The "hits" to one or more database sequences by a queried sequence produced by BLASTN, FASTA, BLASTP or a similar algorithm, align and identify similar portions of sequences. The hits are arranged in order of the degree of similarity and the length of sequence overlap. Hits to a database sequence generally represent an overlap over only a fraction of the sequence length of the queried sequence.

The BLASTN, FASTA, and BLASTP algorithms also produce "Expect" values for alignments. The Expect value (E) indicates the number of hits one can "expect" to see over a certain number of contiguous sequences by chance when searching a database of a certain size. The Expect value is used as a significance threshold for determining whether  
5 the hit to a database, such as the preferred EMBL database, indicates true similarity. For example, an E value of 0.1 assigned to a polynucleotide hit is interpreted as meaning that in a database of the size of the EMBL database, one might expect to see 0.1 matches over the aligned portion of the sequence with a similar score simply by chance. By this criterion, the aligned and matched portions of the polynucleotide sequences then have a  
10 probability of 90% of being the same. For sequences having an E value of 0.01 or less over aligned and matched portions, the probability of finding a match by chance in the EMBL database is 1% or less using the BLASTN or FASTA algorithm.

According to one embodiment, "variant" polynucleotides and polypeptides, with reference to each of the polynucleotides and polypeptides of the present invention,  
15 preferably comprise sequences producing an E value of 0.01 or less when compared to the polynucleotide or polypeptide of the present invention. That is, a variant polynucleotide or polypeptide is any sequence that has at least a 99% probability of being the same as the polynucleotide or polypeptide of the present invention, measured as having an E value of 0.01 or less using the BLASTN, FASTA, or BLASTP algorithms set at parameters  
20 described above. According to a preferred embodiment, a variant polynucleotide is a sequence having the same number or fewer nucleic acids than a polynucleotide of the present invention that has at least a 99% probability of being the same as the polynucleotide of the present invention, measured as having an E value of 0.01 or less using the BLASTN or FASTA algorithms set at parameters described above. Similarly,  
25 according to a preferred embodiment, a variant polypeptide is a sequence having the same number or fewer amino acids than a polypeptide of the present invention that has at least a 99% probability of being the same as a polypeptide of the present invention, measured as having an E value of 0.01 or less using the BLASTP algorithm set at the parameters described above.

30 Alternatively, variant polynucleotides or polypeptides of the present invention comprise a sequence exhibiting at least 40%; more preferably at least 60%; more preferably yet at least 75%; and most preferably at least 90% identity to a polynucleotide or polypeptide of the present invention, determined as described below. The percentage

identity is determined by aligning sequences using one of the BLASTN, FASTA, or BLASTP algorithms, set at the running parameters described above, and identifying the number of identical nucleic or amino acids over the aligned portions; dividing the number of identical nucleic or amino acids by the total number of nucleic or amino acids of the polynucleotide or polypeptide of the present invention; and then multiplying by 100 to determine the percentage identity. For example, a polynucleotide of the present invention having 220 nucleic acids has a hit to a polynucleotide sequence in the EMBL database having 520 nucleic acids over a stretch of 23 nucleotides in the alignment produced by the BLASTN algorithm using the parameters described above. The 23 nucleotide hit includes 21 identical nucleotides, one gap and one different nucleotide. The percentage identity of the polynucleotide of the present invention to the hit in the EMBL library is thus 21/220 times 100, or 9.5%. The polynucleotide sequence in the EMBL database is thus not a variant of a polynucleotide of the present invention.

Alternatively, variant polynucleotides of the present invention hybridize to the polynucleotide sequences recited in SEQ ID NOS: 1-53 and 78-164, or complements, reverse sequences, or reverse complements of those sequences under stringent conditions. As used herein, "stringent conditions" refers to prewashing in a solution of 6X SSC, 0.2% SDS; hybridizing at 65°C, 6X SSC, 0.2% SDS overnight; followed by two washes of 30 minutes each in 1X SSC, 0.1% SDS at 65°C and two washes of 30 minutes each in 0.2X SSC, 0.1% SDS at 65°C.

The present invention also encompasses polynucleotides that differ from the disclosed sequences but that, as a consequence of the discrepancy of the genetic code, encode a polypeptide having similar enzymatic activity as a polypeptide encoded by a polynucleotide of the present invention. Thus, polynucleotides comprising sequences that differ from the polynucleotide sequences recited in SEQ ID NOS: 1-53 and 78-164, or complements, reverse sequences, or reverse complements of those sequences as a result of conservative substitutions are contemplated by and encompassed within the present invention. Additionally, polynucleotides comprising sequences that differ from the polynucleotide sequences recited in SEQ ID NOS: 1-53 and 78-164, or complements, reverse complements, or reverse sequences as a result of deletions and/or insertions totaling less than 10% of the total sequence length are also contemplated by and encompassed within the present invention. Similarly, polypeptides comprising sequences that differ from the polypeptide sequences recited in SEQ ID NOS: 165-304 as a result of

amino acid substitutions, insertions, and/or deletions totaling less than 10% of the total sequence length are contemplated by an encompassed within the present invention, provided the variant polypeptide has activity in an isoprenoid biosynthetic pathway.

The polynucleotides of the present invention may be isolated from various  
5 libraries, or may be synthesized using techniques that are well known in the art. The polynucleotides may be synthesized, for example, using automated oligonucleotide synthesizers (e.g., Beckman Oligo 1000M DNA Synthesizer) to obtain polynucleotide segments of up to 50 or more nucleic acids. A plurality of such polynucleotide segments may then be ligated using standard DNA manipulation techniques that are well known in  
10 the art of molecular biology. One conventional and exemplary polynucleotide synthesis technique involves synthesis of a single stranded polynucleotide segment having, for example, 80 nucleic acids, and hybridizing that segment to a synthesized complementary 85 nucleic acid segment to produce a 5 nucleotide overhang. The next segment may then be synthesized in a similar fashion, with a 5 nucleotide overhang on the opposite strand.  
15 The "sticky" ends ensure proper ligation when the two portions are hybridized. In this way, a complete polynucleotide of the present invention may be synthesized entirely *in vitro*.

Some of the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164 are referred to as "partial" sequences, in that they do not represent the full coding portion of a  
20 gene encoding a naturally occurring polypeptide. The partial polynucleotide sequences disclosed herein may be employed to obtain the corresponding full length genes for various species and organisms by, for example, screening DNA expression libraries using hybridization probes based on the polynucleotides of the present invention, or using PCR amplification with primers based upon the polynucleotides of the present invention. In this  
25 way one can, using methods well known in the art, extend a polynucleotide of the present invention upstream and downstream of the corresponding mRNA, as well as identify the corresponding genomic DNA, including the promoter and enhancer regions, of the complete gene. The present invention thus comprehends isolated polynucleotides comprising a sequence identified in SEQ ID NOS: 1-53 and 78-164, or a variant of one of  
30 the specified sequences, that encode a functional polypeptide, including full length genes. Such extended polynucleotides may have a length of from about 50 to about 4,000 nucleic acids or base pairs, and preferably have a length of less than about 4,000 nucleic acids or base pairs, more preferably yet a length of less than about 3,000 nucleic acids or base

pairs, more preferably yet a length of less than about 2,000 nucleic acids or base pairs. Under some circumstances, extended polynucleotides of the present invention may have a length of less than about 1,800 nucleic acids or base pairs, preferably less than about 1,600 nucleic acids or base pairs, more preferably less than about 1,400 nucleic acids or base  
5 pairs, more preferably yet less than about 1,200 nucleic acids or base pairs, and most preferably less than about 1,000 nucleic acids or base pairs.

Polynucleotides of the present invention also comprehend polynucleotides comprising at least a specified number of contiguous residues ( $x$ -mers) of any of the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164, complements, reverse  
10 sequences, and reverse complements of such sequences, and their variants. Similarly, polypeptides of the present invention comprehend polypeptides comprising at least a specified number of contiguous residues ( $x$ -mers) of any of the polypeptides identified as SEQ ID NOS: 165-304, and their variants. As used herein, the term " $x$ -mer," with  
15 reference to a specific value of " $x$ ," refers to a sequence comprising at least a specified number (" $x$ ") of contiguous residues of any of the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164, or the polypeptides identified as SEQ ID NOS: 165-304. According to preferred embodiments, the value of  $x$  is preferably at least 20; more preferably, at least 40; more preferably yet, at least 60; and most preferably, at least 80. Thus, polynucleotides and polypeptides of the present invention comprise a 20-mer, a 40-  
20 mer, a 60-mer, an 80-mer, a 100-mer, a 120-mer, a 150-mer, a 180-mer, a 220-mer, a 250-mer, or a 300-mer, 400-mer, 500-mer or 600-mer of a polynucleotide or polypeptide identified as SEQ ID NOS: 1-53, and 78-304, and variants thereof.

Polynucleotide probes and primers complementary to and/or corresponding to SEQ ID NOS: 1-53 and 78-164, and variants of those sequences, are also comprehended by the  
25 present invention. Such oligonucleotide probes and primers are substantially complementary to the polynucleotide of interest. As used herein, the term "oligonucleotide" refers to a relatively short segment of a polynucleotide sequence, generally comprising between 6 and 60 nucleotides, and comprehends both probes for use in hybridization assays and primers for use in the amplification of DNA by polymerase  
30 chain reaction.

An oligonucleotide probe or primer is described as "corresponding to" a polynucleotide of the present invention, including one of the sequences set out as SEQ ID NOS: 1-53 and 78-164, or a variant, if the oligonucleotide probe or primer, or its



complement, is contained within one of the sequences set out as SEQ ID NOS: 1-53 and 78-164, or a variant of one of the specified sequences.

Two single stranded sequences are said to be substantially complementary when the nucleotides of one strand, optimally aligned and compared, with the appropriate  
5 nucleotide insertions and/or deletions, pair with at least 80%, preferably at least 90% to 95%, and more preferably at least 98% to 100%, of the nucleotides of the other strand. Alternatively, substantial complementarity exists when a first DNA strand selectively hybridizes to a second DNA strand under stringent hybridization conditions. Stringent hybridization conditions for determining complementarity include salt conditions of less  
10 than about 1 M, more usually less than about 500 mM and preferably less than about 200 mM. Hybridization temperatures may be as low as 5°C, but are generally greater than about 22°C, more preferably greater than about 30°C and most preferably greater than about 37°C. Longer DNA fragments may require higher hybridization temperatures for specific hybridization. Since the stringency of hybridization may be affected by other  
15 factors such as probe composition, presence of organic solvents and extent of base mismatching, the combination of parameters is more important than the absolute measure of any one alone. The DNA from plants or samples or products containing plant material can be either genomic DNA or DNA derived by preparing cDNA from the RNA present in the sample.

20 In addition to DNA-DNA hybridization, DNA-RNA or RNA-RNA hybridization assays are also possible. In the first case, the mRNA from expressed genes would then be detected instead of genomic DNA or cDNA derived from mRNA of the sample. In the second case, RNA probes could be used. In addition, artificial analogs of DNA hybridizing specifically to target sequences could also be used.

25 In specific embodiments, the oligonucleotide probes and/or primers comprise at least about 6 contiguous residues, more preferably at least about 10 contiguous residues, and most preferably at least about 20 contiguous residues complementary to a polynucleotide sequence of the present invention. Probes and primers of the present invention may be from about 8 to 100 base pairs in length or, preferably, from about 10 to  
30 50 base pairs in length or, more preferably, from about 15 to 40 base pairs in length. The probes can be easily selected using procedures well known in the art, taking into account DNA-DNA hybridization stringencies, annealing and melting temperatures, potential for formation of loops and other factors, which are well known in the art. Tools and software

suitable for designing probes, and especially suitable for designing PCR primers, are available on the Internet, for example, at URL <http://www.horizonpress.com/pcr/>. Preferred techniques for designing PCR primers are also disclosed in Dieffenbach CW and Dyksler GS, *PCR primer: a laboratory manual*. CSHL Press: Cold Spring Harbor, NY, 5 1995.

A plurality of oligonucleotide probes or primers corresponding to a polynucleotide of the present invention may be provided in a kit form. Such kits generally comprise multiple DNA or oligonucleotide probes, each probe being specific for a polynucleotide sequence. Kits of the present invention may comprise one or more probes or primers 10 corresponding to a polynucleotide of the present invention, including a polynucleotide sequence identified in SEQ ID NOS: 1-53 and 78-164.

In one embodiment useful for high-throughput assays, the oligonucleotide probe kits of the present invention comprise multiple probes in an array format, wherein each probe is immobilized in a predefined, spatially addressable location on the surface of a 15 solid substrate. Array formats which may be usefully employed in the present invention are disclosed, for example, in U.S. Patent Nos. 5,412,087 and 5,545,531; and PCT Publication No. WO 95/00530, the disclosures of which are hereby incorporated by reference.

Probes, preferably in the form of an array, may be employed to screen for 20 differences in organisms or samples or products containing genetic material using high throughput screening techniques that are well known in the art. The significance of using probes in high-throughput screening systems is apparent for applications such as plant breeding and quality control operations in which there is a need to identify large numbers of seed lots and plant seedlings, to examine samples or products for unwanted plant 25 materials, to identify plants or samples or products containing plant material for quarantine purposes, etc., or to ascertain the true origin of plants or samples or products containing plant material. Screening for the presence or absence of polynucleotides of the present invention used as identifiers for tagging plants is valuable for later detecting the amount of gene flow in plant breeding, introgression of genes via dispersed pollen, etc.

30 In this manner, oligonucleotide probe kits of the present invention may be employed to examine the presence/absence (or relative amounts in case of mixtures) of polynucleotides in different samples or products containing different materials rapidly and in a cost-effective manner. Examples of plant species that may be examined using the

present invention, include forestry species, such as pine and eucalyptus species, other tree species, and agricultural and horticultural plants.

Another aspect of the present invention involves collections of a plurality of polynucleotides of the present invention. A collection of a plurality of the polynucleotides  
5 of the present invention, particularly the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164, and variants thereof, may be recorded and/or stored on a storage medium and subsequently accessed for purposes of analysis, comparison, etc. Suitable storage media include magnetic media such as magnetic diskettes, magnetic tapes, CD-ROM storage media, optical storage media, and the like. Suitable storage media and methods for  
10 recording and storing information, as well as accessing information such as polynucleotide sequences recorded on such media, are well known in the art. The polynucleotide information stored on the storage medium is preferably computer-readable and may be used for analysis and comparison of the polynucleotide information.

According to one embodiment, the storage medium includes a collection of at least  
15 4, preferably at least 10, more preferably at least 15, and most preferably at least 20 of the polynucleotides of the present invention, preferably the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164, and variants of those polynucleotides.

For applications where modulation of a polypeptide involved with isoprenoid biosynthesis and/or isoprenoid metabolism is desired, an open reading frame may be  
20 inserted into a genetic construct in a sense or antisense orientation, such that transformation of a target plant with the genetic construct produces a change in the expression level of the polypeptide compared to the expression in a wild-type organism. Transformation with a genetic construct comprising an open reading frame in a sense orientation will generally result in modulation of expression of the selected gene, while  
25 transformation with a genetic construct comprising an open reading frame in an antisense orientation generally produces reduced expression of the selected gene. A population of plants transformed with a genetic construct comprising an open reading frame of the present invention in either a sense or antisense orientation may be screened for increased or reduced expression of the gene in question using techniques well known to those of  
30 skill in the art, and plants having the desired phenotypes may thus be isolated.

Alternatively, expression of a gene involved in the biosynthesis of isoprenoids may be inhibited by inserting a portion of an open reading frame of the present invention, in either sense or antisense orientation, in the genetic construct. Such portions need not be

full-length but preferably comprise at least 25, and more preferably, at least 50 residues of polynucleotide of the present invention. A much longer portion, or even the full length polynucleotide corresponding to the complete open reading frame, may be employed. The portion of the open reading frame does not need to be precisely the same as the  
5 endogenous sequence, provided that there is sufficient sequence similarity to achieve inhibition of the target gene. Thus a sequence derived from one species may be used to inhibit expression of a gene in a different species.

According to another embodiment, the genetic constructs of the present invention comprise a polynucleotide including a non-coding region of a gene coding for a polypeptide encoded by a polynucleotide of the present invention, or a polynucleotide  
10 complementary to such a non-coding region. Examples of non-coding regions which may be usefully employed in such constructs include introns and 5'-non-coding leader sequences. Transformation of a target plant with such a genetic construct may lead to a reduction in the amount of an isoprenoid compound synthesized by the plant by the  
15 process of cosuppression, in a manner similar to that discussed, for example, by Napoli *et al.*, *Plant Cell* 2:279-290, 1990 and de Carvalho Niebel *et al.*, *Plant Cell* 7:347-358, 1995.

Alternatively, regulation may be achieved by inserting appropriate sequences or subsequences (e.g. DNA or RNA) in ribozyme constructs (McIntyre CL and Manners JM, *Transgenic Res.* 5(4):257-262, 1996). Ribozymes are synthetic RNA molecules that  
20 comprise a hybridizing region complementary to two regions, each of which comprises at least 5 contiguous nucleotides in a mRNA molecule encoded by one of the inventive polynucleotides. Ribozymes possess highly specific endonuclease activity, which autocatalytically cleaves the mRNA.

The genetic constructs of the present invention further comprise a gene promoter  
25 sequence and a gene termination sequence, operably linked to the polynucleotide to be transcribed, which control expression of the polypeptide. The gene promoter sequence is generally positioned at the 5' end of the polynucleotide to be transcribed, and is employed to initiate transcription of the polynucleotide. Gene promoter sequences are generally found in the 5' non-coding region of a gene but they may exist downstream of the open  
30 reading frame or in introns (Luehrsen KR, *Mol. Gen. Genet.* 225:81-93, 1991); or in the coding region, as for example in a plant defence gene (Douglas *et al.*, *EMBO J.* 10:1767-1775, 1991). When the construct includes an open reading frame in a sense orientation, the gene promoter sequence also initiates translation of the open reading frame. For

genetic constructs comprising either an open reading frame in an antisense orientation or a non-coding region, the gene promoter sequence consists only of a transcription initiation site having a RNA polymerase binding site.

Numerous gene promoter sequences that may be usefully employed in the genetic  
5 constructs of the present invention are well known in the art. The gene promoter sequence, and also the gene termination sequence, may be endogenous to the target plant host or may be exogenous, provided the promoter is functional in the target host. For example, the promoter and termination sequences may be from other plant species, plant viruses, bacterial plasmids and the like. Preferably, gene promoter and termination  
10 sequences are common to those of the polynucleotide being introduced.

Factors influencing the choice of promoter include the desired tissue specificity of the construct, and the timing of transcription and translation. For example, constitutive promoters, such as the 35S Cauliflower Mosaic Virus (CaMV 35S) promoter with or without enhancers, such as the Kozak sequence or the Omega enhancer, and  
15 *Agrobacterium tumefaciens* nopaline synthase terminator, may be usefully employed in the present invention. Use of a tissue specific promoter will result in production of the desired sense or antisense RNA only in the tissue of interest. With genetic constructs employing inducible gene promoter sequences, the rate of RNA polymerase binding and initiation can be modulated by external stimuli, such as light, heat, anaerobic stress, alteration in nutrient  
20 conditions and the like. Temporally regulated promoters can be employed to effect modulation of the rate of RNA polymerase binding and initiation at a specific time during development of a transformed cell. Preferably, the original promoters from the enzyme gene in question, or promoters from a specific tissue-targeted gene in the organism to be transformed, such as eucalyptus or pine are used. Other examples of gene promoters  
25 which may be usefully employed in the present invention include mannopine synthase (mas), octopine synthase (ocs) and those reviewed by Chua *et al.*, *Science* 244:174-181, 1989.

The gene termination sequence, which is located 3' to the polynucleotide to be transcribed, may come from the same gene as the gene promoter sequence or may be from  
30 a different gene. Many gene termination sequences known in the art may be usefully employed in the present invention, such as the 3' end of the *Agrobacterium tumefaciens* nopaline synthase gene. However, preferred gene terminator sequences are those from the original enzyme gene or from the target species to be transformed.

The genetic constructs of the present invention may also contain a selection marker that is effective in target cells, such as plant cells, to allow for the detection of transformed cells containing the inventive construct. Such markers, which are well known in the art, typically confer resistance to one or more toxins. One example of such a marker is the NPTII gene whose expression results in resistance to kanamycin or hygromycin, antibiotics which are usually toxic to plant cells at a moderate concentration (Rogers et al. in Weissbach A and Weissbach H, eds., *Methods for Plant Molecular Biology*, Academic Press Inc.: San Diego, CA, 1988). Transformed cells can thus be identified by their ability to grow in media containing the antibiotic in question. Alternatively, the presence of the desired construct in transformed cells can be determined by means of other techniques well known in the art, such as Southern and Western blots. A transcription initiation site may additionally included in the genetic construct when the sequence to be transcribed lacks such a site.

Techniques for operatively linking the components of the genetic constructs of the present invention are well known in the art and include the use of synthetic linkers containing one or more restriction endonuclease sites as described, for example, by Sambrook et al., *Molecular cloning: a laboratory manual*, CSHL Press: Cold Spring Harbor, NY, 1989. The DNA construct of the present invention may be linked to a vector having at least one replication system, for example *E. coli*, whereby after each manipulation, the resulting construct can be cloned and sequenced and the correctness of the manipulation determined.

The genetic constructs of the present invention may be used to transform a variety of target organisms such as plants, both monocotyledonous (e.g., grasses, corn, grains, oat, wheat and barley); dicotyledonous (e.g., *Arabidopsis*, tobacco, legumes, alfalfa, oaks, eucalyptus, maple); gymnosperms (e.g., Scots pine (Aronen, *Finnish Forest Res. Papers*, Vol. 595, 1996); white spruce (Ellis et al., *Biotechnology* 11: 84-89, 1993); and larch (Huang et al., *In Vitro Cell* 27:201-207, 1991). In a preferred embodiment, the inventive DNA constructs are employed to transform woody plants, herein defined as a tree or shrub whose stem lives for a number of years and increases in diameter each year by the addition of woody tissue. Preferably the target plant is selected from the group consisting of eucalyptus and pine species, most preferably from the group consisting of *Eucalyptus grandis* and *Pinus radiata*. Other species which may be usefully transformed with the DNA constructs of the present invention include, but are not limited to: Pines, such as

*Pinus banksiana*, *Pinus brutia*, *Pinus caribaea*, *Pinus clausa*, *Pinus contorta*, *Pinus coulteri*, *Pinus echinata*, *Pinus eldarica*, *Pinus ellioti*, *Pinus jeffreyi*, *Pinus lambertiana*, *Pinus monticola*, *Pinus nigra*, *Pinus palustris*, *Pinus pinaster*, *Pinus ponderosa*, *Pinus resinosa*, *Pinus rigida*, *Pinus serotina*, *Pinus strobus*, *Pinus sylvestris*, *Pinus taeda*, *Pinus virginiana*; other gymnosperm, such as *Abies amabilis*, *Abies balsamea*, *Abies concolor*, *Abies grandis*, *Abies lasiocarpa*, *Abies magnifica*, *Abies procera*, *Chamaecyparis lawsoniana*, *Chamaecyparis nootkatensis*, *Chamaecyparis thyoides*, *Huniperus virginiana*, *Larix decidua*, *Larix laricina*, *Larix leptolepis*, *Larix occidentalis*, *Larix siberica*, *Libocedrus decurrens*, *Picea abies*, *Picea engelmanni*, *Picea glauca*, *Picea mariana*, *Picea pungens*, *Picea rubens*, *Picea sitchensis*, *Pseudotsuga menziesii*, *Sequoia gigantea*, *Sequoia sempervirens*, *Taxodium distichum*, *Tsuga canadensis*, *Tsuga heterophylla*, *Tsuga mertensiana*, *Thuja occidentalis*, *Thuja plicata*; and Eucalypts, such as *Eucalyptus alba*, *Eucalyptus bancroftii*, *Eucalyptus botyroides*, *Eucalyptus bridgesiana*, *Eucalyptus calophylla*, *Eucalyptus camaldulensis*, *Eucalyptus citriodora*, *Eucalyptus cladocalyx*, *Eucalyptus coccifera*, *Eucalyptus curtisii*, *Eucalyptus dalrympleana*, *Eucalyptus deglupta*, *Eucalyptus delagatensis*, *Eucalyptus diversicolor*, *Eucalyptus dunnii*, *Eucalyptus ficifolia*, *Eucalyptus globulus*, *Eucalyptus gomphocephala*, *Eucalyptus gunnii*, *Eucalyptus henryi*, *Eucalyptus laevopinea*, *Eucalyptus macarthurii*, *Eucalyptus macrorhyncha*, *Eucalyptus maculata*, *Eucalyptus marginata*, *Eucalyptus megacarpa*, *Eucalyptus melliodora*, *Eucalyptus nicholii*, *Eucalyptus nitens*, *Eucalyptus nova-anglica*, *Eucalyptus obliqua*, *Eucalyptus obtusiflora*, *Eucalyptus oreades*, *Eucalyptus pauciflora*, *Eucalyptus polybractea*, *Eucalyptus regnans*, *Eucalyptus resinifera*, *Eucalyptus robusta*, *Eucalyptus rudis*, *Eucalyptus saligna*, *Eucalyptus sideroxylon*, *Eucalyptus stuartiana*, *Eucalyptus tereticornis*, *Eucalyptus torelliana*, *Eucalyptus urnigera*, *Eucalyptus urophylla*, *Eucalyptus viminalis*, *Eucalyptus viridis*, *Eucalyptus wandoo*, *Eucalyptus youmanni*.

Techniques for stably incorporating genetic constructs into the genome of target plants are well known in the art and include *Agrobacterium tumefaciens* mediated introduction, electroporation, protoplast fusion, injection into reproductive organs, injection into immature embryos, high velocity projectile introduction, and the like. The choice of technique will depend upon the target plant to be transformed. For example, dicotyledonous plants and certain monocots and gymnosperms may be transformed by *Agrobacterium* Ti plasmid technology, as described, for example by Bevan, *Nucleic Acid Res.* 12:8711-8721, 1984. Targets for the introduction of the genetic constructs of the

present invention include tissues, such as leaf tissue, disseminated cells, protoplasts, seeds, embryos, meristematic regions; cotyledons, hypocotyls, and the like. The preferred method for transforming eucalyptus and pine is a biolistic method using pollen (*see, for example, Aronen, Finnish Forest Res. Papers 595:53, 1996*) or easily regenerable  
5 embryonic tissues.

Once the cells are transformed, cells having the inventive genetic construct incorporated in their genome may be selected by means of a marker, such as the kanamycin resistance marker discussed above. Transgenic cells may then be cultured in an appropriate medium to regenerate whole plants, using techniques well known in the art.  
10 In the case of protoplasts, the cell wall is allowed to reform under appropriate osmotic conditions. In the case of seeds or embryos, an appropriate germination or callus initiation medium is employed. For explants, an appropriate regeneration medium is used. Regeneration of plants is well established for many species. For a review of regeneration of forest trees, *see Dunstan et al., in Thorpe TA, ed., In vitro embryogenesis of plants,*  
15 *Current Plant Science and Biotechnology in Agriculture, 20(12):471-540, 1995.* Specific protocols for the regeneration of spruce are discussed by Roberts., Somatic embryogenesis of spruce," in Redenbaugh K, ed., *Synseed: applications of synthetic seed to crop improvement*, CRC Press: Ch. 23, pp. 427-449, 1993. The resulting transformed plants may be reproduced sexually or asexually, using methods well known in the art, to give  
20 successive generations of transgenic plants.

As discussed above, the production of RNA in target plant cells can be controlled by choice of the promoter sequence, or by selecting the number of functional copies or the site of integration of the polynucleotides incorporated into the genome of the target plant host. A target plant may be transformed with more than one genetic constructs of the  
25 present invention, thereby modulating the activity of more than one isoprenoid metabolism enzyme, affecting enzyme activity in more than one tissue, or affecting enzyme activity at more than one expression time. Similarly, a genetic construct may be assembled containing more than one open reading frame coding for an enzyme encoded by a polynucleotide of the present invention or more than one non-coding region of a gene  
30 coding for such an enzyme. The polynucleotides of the present inventive may also be employed in combination with other known sequences encoding enzymes involved in the synthesis of isoprenoids.



Additionally, the polynucleotides of the present invention have particular application for use as non-disruptive tags for marking organisms, particularly plants. Genetic constructs comprising polynucleotides of the present invention may be stably introduced into an organism as heterologous, non-functional, non-disruptive tags. It is  
5 then possible to identify the origin or source of the organism at a later date by determining the presence or absence of the tag(s) in a sample of material. Organisms other than plants may also be tagged with the polynucleotides of the present invention, including commercially valuable animals, fish, bacteria and yeasts.

Detection of the tag(s) may be accomplished using a variety of conventional  
10 techniques, and will generally involve the use of nucleic acid probes. Sensitivity in assaying the presence of probe can be usefully increased by using branched oligonucleotides, as described by Horn *et al.*, *Nucleic Acids Res.* 25(23):4842-4849, 1997), enabling detection of as few as 50 DNA molecules in the sample.

The following examples are offered by way of illustration and not by way of  
15 limitation.

#### Example 1

##### Isolation and Characterization of cDNA Clones from *Pinus radiata* and *Eucalyptus grandis*

*Pinus radiata* and *Eucalyptus grandis* cDNA expression libraries were constructed  
20 and screened as follows. mRNA was extracted from the plant tissue using the protocol of Chang *et al.*, *Plant Molecular Biology Reporter* 11:113-116, 1993 with minor modifications. Specifically, samples were dissolved in CPC-RNAXB (100 mM Tris-Cl, pH 8.0; 25 mM EDTA; 2.0 M NaCl; 2%CTAB; 2% PVP and 0.05% Spermidine\*3HCl) and extracted with chloroform:isoamyl alcohol, 24:1. mRNA was precipitated with  
25 ethanol and the total RNA preparate was purified using a Poly(A) Quik mRNA Isolation Kit (Stratagene, La Jolla, CA). A cDNA expression library was constructed from the purified mRNA by reverse transcriptase synthesis followed by insertion of the resulting cDNA clones in Lambda ZAP using a ZAP Express cDNA Synthesis Kit (Stratagene), according to the manufacturer's protocol. The resulting cDNAs were packaged using a  
30 Gigapack II Packaging Extract (Stratagene) employing 1 µl of sample DNA from the 5 µl ligation mix. Mass excision of the library was done using XL1-Blue MRF' cells and XL0LR cells (Stratagene) with ExAssist helper phage (Stratagene). The excised phagemids were diluted with NZY broth (Gibco BRL, Gaithersburg, MD) and plated out

onto LB-kanamycin agar plates containing X-gal and isopropylthio-beta-galactoside (IPTG).

Of the colonies plated and picked for DNA miniprep, the large majority contained an insert suitable for sequencing. Positive colonies were cultured in NZY broth with kanamycin and cDNA was purified by means of REAL DNA minipreps (Qiagen, Venlo, The Netherlands). Agarose gel at 1% was used to screen sequencing templates for chromosomal contamination. Dye terminator sequences were prepared using a Biomek 2000 robot (Beckman Coulter Inc, Fullerton CA for liquid handling and DNA amplification using a 9700 PCR machine (Perkin Elmer/Applied Biosystems, Foster City, CA) according to the manufacturer's protocol.

Polynucleotides for positive clones were obtained using a Perkin Elmer/Applied Biosystems Division Prism 377 sequencer. cDNA clones were sequenced first from the 5' end and, in some cases, also from the 3' end. For some clones, internal sequences were obtained using subcloned fragments. Subcloning was performed using standard procedures of restriction mapping and subcloning to pBluescript II SK+ vector and other standard sequencing vectors.

The determined cDNA sequences, including the polynucleotides of the present invention, were compared to and aligned with known sequences in the. Specifically, the polynucleotides identified in SEQ ID NOS. 1-53 were compared to polynucleotides in the EMBL database EMBL as of the end of August, 1998 using the BLASTN algorithm Version 2.0.4 [Feb-24-1998] set to the following running: Unix running command: blastall -p blastn -d embldb -e 10 -G 0 -E 0 -r 1 -v 30 -b 30 -i queryseq -o results. The polynucleotides identified in SEQ ID NOS: 78-164 were compared to polynucleotides in the EMBL database EMBL as of the end of May, 1999 using BLASTN algorithm Version 2.0.6 [Sep-16-1998], set to the following running parameters: Unix running command: blastall -p blastn -d embldb -e 10 -G 0 -E 0 -r 1 -v 30 -b 30 -i queryseq -o results. Multiple alignments of redundant sequences were used to build up reliable consensus sequences. Based on similarity to known sequences from other plant species, the isolated polynucleotides of the present invention identified as SEQ ID NOS. 1-53 and 78-164 were putatively identified as encoding polypeptides having similarity to the polypeptides shown above in Table 1.

The isolated cDNA sequences were compared to sequences in the EMBL DNA database using the computer algorithm BLASTN. The corresponding predicted

polypeptide sequences were determined and were compared to sequences in the SwissProt database using the computer algorithm BLASTP. Comparisons of DNA sequences provided in SEQ ID NOS: 78-164, to sequences in the EMBL DNA database (using BLASTN) and amino acid sequences provided in SEQ ID NOS: 165-304 to sequences in the SwissProt database (using BLASTP) were made as of May, 1999. Analysis of six-frame translations of the polynucleotides of SEQ ID NOS: 78-164, were also compared to and aligned with the six-frame translations of polynucleotides in the EMBL database using the TBLASTX program.

10 BLASTN Polynucleotide Analysis

The cDNA sequences of SEQ ID NOS: 1, 2, 4-6, 8-12, 15, 19, 21-23, 27-33, 35, 37-42, 44, 46-52, 78-80, 82, 83, 86, 89-92, 96-100, 104-113, 115, 117, 120, 122-130, 132-136, 138-158, 160, 163 and 164, were determined to have less than 40% identity, determined as described above, to sequences in the EMBL database using the computer algorithm BLASTN, as described above. The cDNA sequences of SEQ ID NOS: 3, 7, 14, 18, 20, 25, 34, 36, 53, 84, 85, 87, 88, 101, 114, 116, 118, 119, 131, 137, 159, 161 and 162 were determined to have less than 60% identity, determined as described above, to sequences in the EMBL database using BLASTN, as described above. The cDNA sequences of SEQ ID NOS: 16, 17, 26, 43, 45, 93, 94 and 121, were determined to have less than 75% identity, determined as described above, to sequences in the EMBL database using BLASTN, as described above. The cDNA sequences of SEQ ID NOS: 13, 24, 95, 102 and 103 were determined to have less than 90% identity, determined as described above, to sequences in the EMBL database using BLASTN, as described above.

25 BLASTP Amino Acid Analysis

The predicted amino acid sequences of SEQ ID NOS: 194-200, 202, 216, 223, 230, 235, 239, 240, 243, 250, 255, 259, 260, 263, 270, 272, 274, 278, 291, 292, 293, 296, 303 and 304 were determined to have less than 50% identity, determined as described above, to sequences in the SwissProt database using the BLASTP computer algorithm as described above. The predicted amino acid sequences of SEQ ID NOS: 166, 168-177, 179, 183-188, 192, 203-205, 207, 209-213, 218, 219, 221, 224, 225, 227-229, 231, 232, 234, 237, 242, 244, 245, 251, 253, 262, 267, 268, 269, 273, 276, 277, 279, 281, 282, 284, 286, 289, 290, 294, 295, 297, 298, 299, 300, 301 and 302 were determined to have less

than 75% identity, determined as described above, to sequences in the SwissProt database using the computer algorithm BLASTP, as described above. The predicted amino acid sequences of SEQ ID NOS: 165, 167, 178, 182, 189-191, 193, 201, 206, 208, 214, 215, 217, 220, 222, 226, 233, 238, 241, 246-250, 254, 256, 257, 258, 261, 264, 265, 266, 275, 5 280, 283, 285 and 288 were determined to have less than 90% identity, determined as described above, to sequences in the SwissProt database using the computer algorithm BLASTP, as described above. The predicted amino acid sequences of SEQ ID NOS: 180, 181 and 271, were determined to have less than 95% identity, determined as described above, to sequences in the SwissProt database using the computer algorithm BLASTP, as described above. 10 described above.

#### TBLASTX Analysis

The six-frame translations of the polynucleotide sequences of SEQ ID NOS: 78-164 were compared to and aligned with six-frame translations of polynucleotides in the EMBL database using the TBLASTX program version 2.0.6 [Sept-16-1998] set to the following running parameters: Unix running command: blastall -p blastn -d embldb -e 10 -G 0 -E 0 -v 30 -b 30 -i queryseq -o results. The translations of the polynucleotides of SEQ ID NOS: 82, 83, 90, 107-113, 115, 120, 122, 124-126, 129, 134-136, 142-144, 146-149, 152, 153, 155-158 and 164, were determined to have less than 50% identity, 15 determined as described above, to translations of polynucleotides in the EMBL database using the computer algorithm TBLASTX. The translations of the polynucleotides of SEQ ID NOS: 79, 81, 84-89, 91, 92, 96-101, 103, 105, 114, 116-118, 123, 131, 132, 137-141, 145, 150, 154 and 160-162, were determined to have less than 75% identity, determined as described above, to translations of polynucleotides in the EMBL database using the computer algorithm TBLASTX. The translations of the polynucleotide sequences of SEQ 25 ID NOS: 78, 80, 93, 95, 102, 104, 106, 119, 121, 127, 128, 130, 133, 151, 159 and 163, were determined to have less than 90% identity, determined as described above, to translations of polynucleotides in the EMBL database using the computer algorithm TBLASTX. The translations of the polynucleotide sequence of SEQ ID NO: 94 was 30 determined to have less than 95% identity, determined as described above, to translations of polynucleotides in the EMBL database using the computer algorithm TBLASTX.

## Example 2

### Use of an O-methyltransferase (OMT) Gene to Modify Lignin Biosynthesis

#### 5    Transformation of tobacco plants with a *Pinus radiata* OMT gene

- Genetic constructs comprising sense and anti-sense nucleotides containing a polynucleotide comprising the coding region of the enzyme O-methyltransferase (OMT) (SEQ ID NO: 54) from *Pinus radiata* were constructed and inserted into *Agrobacterium tumefaciens* by direct transformation using published methods (An *et al.*, "Binary  
10    vectors," in Gelvin SB and Schilperoort RA, eds., *Plant Molecular Biology Manual*, Kluwer Academic Publishers: Dordrecht, 1988). General methods for plant transformation are described in Horsch *et al.*, *Science* 227:1229-1231, 1985. The constructs of sense DNA were made by first cloning the PBK-CMV cDNA inserts into pART7 vectors. The pART7 vectors were then cut by restriction endonuclease *NotI* to remove the 35S-Insert-  
15    OCS 3'UTR construct for cloning into the plant expression vector pART27 (Gleave A, *Plant Mol. Biol.* 20:1203-1207, 1992). The presence and integrity of the transgenic constructs were verified by restriction digestion and DNA sequencing.

Tobacco (*Nicotiana tabacum* cv. Samsun) leaf sections were transformed with the sense and anti-sense OMT constructs using the method of Horsch *et al.*, *Science*  
20    227:1229-1231, 1985. Five independent transformed plant lines were established for the sense construct and eight independent transformed plant lines were established for the anti-sense construct for OMT. Transformed plants containing the appropriate gene construct were verified using Southern blot experiments. A "+" in the column labeled "Southern" in Table 2 below indicates that the transformed plant lines were confirmed as  
25    independent transformed lines.

#### Expression of *Pinus* OMT in transformed plants

Total RNA was isolated from each independent transformed plant line created with the OMT sense and anti-sense constructs. The RNA samples were analyzed in Northern  
30    blot experiments to determine the level of expression of the transgene in each transformed line.

The data shown in the column labeled "Northern" in Table 1 shows that the transformed plant lines containing the sense and anti-sense constructs for OMT all exhibited high levels of expression, relative to the background on the Northern blots.

OMT expression in sense plant line number 2 was not measured because the RNA sample showed signs of degradation. There was no detectable hybridization to RNA samples from empty vector-transformed control plants.

5 Modulation of OMT enzyme activity in transformed plants

The total activity of OMT enzyme, encoded by the *Pinus* OMT gene and by the endogenous tobacco OMT gene, was analyzed for each transformed plant line created with the OMT sense and anti-sense constructs. Crude protein extracts were prepared from each transformed plant and assayed using the method of Zhang *et al.*, *Plant Physiol.* 113:65-74,  
10 1997. The data contained in the column labeled "Enzyme" in Table 2 shows that the transformed plant lines containing the OMT sense construct generally had elevated OMT enzyme activity, with a maximum of 199%, whereas the transformed plant lines containing the OMT anti-sense construct generally had reduced OMT enzyme activity, with a minimum of 35%, relative to empty vector-transformed control plants. OMT  
15 enzyme activity was not estimated in sense plant line number 3.

Effects of *Pinus* OMT on lignin concentration in transformed plants

OMT is an enzyme involved in the biosynthesis of lignin. The concentration of lignin in the transformed tobacco plants was determined using the well-established  
20 procedure of thioglycolic acid extraction (Freudenberg *et al.*, *Constitution and Biosynthesis of Lignin*, Springer-Verlag: Berlin, 1968). Briefly, whole tobacco plants, of an average age of 38 days, were frozen in liquid nitrogen and ground to a fine powder in a mortar and pestle. 100 mg of frozen powder from one empty vector-transformed control plant line, the five independent transformed plant lines containing the sense construct for  
25 OMT and the eight independent transformed plant lines containing the anti-sense construct for OMT were extracted individually with methanol, followed by 10% thioglycolic acid and finally dissolved in 1 M NaOH. The final extracts were assayed for absorbance at 280 nm. The data shown in the column labeled "TGA" in Table 2 shows that the transformed plant lines containing the sense and the anti-sense OMT gene constructs all exhibited  
30 significantly decreased levels of lignin, relative to the empty vector-transformed control plant lines.

TABLE 2

	plant line	transgene	orientation	Southern	Northern	Enzyme	TGA
5	1	control	na	+	blank	100	104
	1	OMT	sense	+	2.9E+6	86	55
	2	OMT	sense	+	na	162	58
	3	OMT	sense	+	4.1E+6	na	63
	4	OMT	sense	+	2.3E+6	142	66
10	5	OMT	sense	+	3.6E+5	199	75
	1	OMT	anti-sense	+	1.6E+4	189	66
	2	OMT	anti-sense	+	5.7E+3	35	70
	3	OMT	anti-sense	+	8.0E+3	105	73
	4	OMT	anti-sense	+	1.4E+4	109	74
15	5	OMT	anti-sense	+	2.5E+4	87	78
	6	OMT	anti-sense	+	2.5E+4	58	84
	7	OMT	anti-sense	+	2.5E+4	97	92
	8	OMT	anti-sense	+	1.1E+4	151	94

20 These data clearly demonstrate that polynucleotides identified from isolated cDNA obtained as in Example 1 and encoding polypeptides, may be assembled in DNA constructs and used to transform plants. The data furthermore demonstrates that transformed plants comprising genetic constructs exhibit varied levels of such enzyme expression and activity, and that the modulation of the metabolism of such an enzyme, manipulated by either sense or anti-sense expression of a gene encoding the enzyme, such as OMT, affects end product concentrations, such as the lignin concentration in the transformed plants.

### Example 3

#### Use of a 4-Coumarate:CoA ligase (4CL) Gene to Modify Lignin Biosynthesis

##### Transformation of tobacco plants with a *Pinus radiata* 4CL gene

35 Sense and anti-sense constructs containing a DNA sequence including the coding region of 4CL (SEQ ID NO: 55) from *Pinus radiata* were inserted into *Agrobacterium tumefaciens* LBA4301 by direct transformation as described above in Example 2. The presence and integrity of the transgenic constructs were verified by restriction digestion and DNA sequencing.

40 Tobacco (*Nicotiana tabacum* cv. Samsun) leaf sections were transformed as described above. Five independent transformed plant lines were established for the sense

construct and eight independent transformed plant lines were established for the anti-sense construct for 4CL. Transformed plants containing the appropriate lignin gene construct were verified using Southern blot experiments. A "+" in the column labeled "Southern" in Table 3 indicates that the transformed plant lines listed were confirmed as independent  
5 transformed lines.

#### Expression of *Pinus* 4CL in transformed plants

Total RNA was isolated from each independent transformed plant line created with the 4CL sense and anti-sense constructs. The RNA samples were analyzed in Northern  
10 blot experiments to determine the level of expression of the transgene in each transformed line. The data shown in the column labeled "Northern" in Table 3 below shows that the transformed plant lines containing the sense and anti-sense constructs for 4CL all exhibit high levels of expression, relative to the background on the Northern blots. 4CL expression in anti-sense plant line number 1 was not measured because the RNA was not  
15 available at the time of the experiment. There was no detectable hybridization to RNA samples from empty vector-transformed control plants.

#### Modulation of 4CL enzyme activity in transformed plants

The total activity of 4CL enzyme, encoded by the *Pinus* 4CL gene and by the  
20 endogenous tobacco 4CL gene in transformed tobacco plants, was analyzed for each transformed plant line created with the 4CL sense and anti-sense constructs. Crude protein extracts were prepared from each transformed plant and assayed using the method of Zhang *et al.*, *Plant Physiol.* 113:65-74, 1997. The data contained in the column labeled "Enzyme" in Table 2 shows that the transformed plant lines containing the 4CL sense  
25 construct had elevated 4CL enzyme activity, with a maximum of 258%, and the transformed plant lines containing the 4CL anti-sense construct had reduced 4CL enzyme activity, with a minimum of 59%, relative to empty vector-transformed control plants.

#### Effects of *Pinus* 4CL on lignin concentration in transformed plants

30 The concentration of lignin in samples of transformed plant material was determined as described in Example 2. The data shown in the column labeled "TGA" in Table 3, below, shows that the transformed plant lines containing the sense and the anti-sense 4CL gene constructs all exhibited significantly decreased levels of lignin, relative to



the empty vector-transformed control plant lines. These data demonstrate that the polynucleotides identified from isolated cDNA as obtained in Example 1 may be assembled into DNA constructs and used to transform plants. Transformed plants comprising such genetic constructs exhibit modified levels of enzyme expression and activity. The metabolism of the biosynthetic pathway involving the enzyme is also affected.

TABLE 3

	plant line	transgene	orientation	Southern	Northern	Enzyme	TGA
10	1	control	na	+	blank	100	92
	2	control	na	+	blank	100	104
	1	4CL	sense	+	2.3E+4	169	64
15	2	4CL	sense	+	4.5E+4	258	73
	3	4CL	sense	+	3.1E+4	174	77
	4	4CL	sense	+	1.7E+4	164	80
	5	4CL	sense	+	1.6E+4	184	92
	1	4CL	anti-sense	+	na	59	75
20	2	4CL	anti-sense	+	1.0E+4	70	75
	3	4CL	anti-sense	+	9.6E+3	81	80
	4	4CL	anti-sense	+	1.2E+4	90	83
	5	4CL	anti-sense	+	4.7E+3	101	88
	6	4CL	anti-sense	+	3.9E+3	116	89
25	7	4CL	anti-sense	+	1.8E+3	125	94
	8	4CL	anti-sense	+	1.7E+4	106	97

Example 430 Transformation of Tobacco using Lignin Biosynthetic Genes

Sense and anti-sense constructs containing DNA sequences including the coding regions of coumarate 3-hydroxylase (C3H) (SEQ ID NO: 56), ferulate-5-hydroxylase (F5H) (SEQ ID NO: 57), cinnamoyl-CoA reductase (CCR) (SEQ ID NO: 58) and coniferyl glycosyl transferase (CGT) (SEQ ID NO: 59) from *Eucalyptus grandis*, and phenylalanine ammonia-lyase (PAL) (SEQ ID NOS: 60 and 61), cinnamate 4-hydroxylase (C4H) (SEQ ID NOS: 62 and 63), phenolase (PNL) (SEQ ID NO: 64) and laccase (LAC) (SEQ ID NO: 65) from *Pinus radiata* were inserted into *Agrobacterium tumefaciens* LBA4301 by direct transformation as described above. The presence and

integrity of the transgenic constructs were verified by restriction digestion and DNA sequencing.

5 Tobacco (*Nicotiana tabacum* cv. Samsun) leaf sections were transformed as described in Example 2. Up to twelve independent transformed plant lines were established for each sense construct and each anti-sense construct listed in the preceding paragraph. Transformed plants containing the appropriate lignin gene construct were verified using Southern blot experiments. All of the transformed plant lines analyzed were confirmed as independent transformed lines. This demonstrates that transgenic plants with an expressed novel gene can be made, starting the whole process from an isolated cDNA  
10 obtained as in Example 1.

### Example 5

#### Manipulation of Lignin Content in Transformed Plants

15

##### Determination of transgene expression by Northern blot experiments

Total RNA was isolated from each independent transformed plant line described in Example 4. The RNA samples were analyzed in Northern blot experiments to determine  
20 the level of expression of the transgene in each transformed line. The column labeled "Northern" in Table 4 shows the level of transgene expression for all plant lines assayed, relative to the background on the Northern blots. There was no detectable hybridization to RNA samples from empty vector-transformed control plants.

##### 25 Determination of lignin concentration in transformed plants

The concentration of lignin in empty vector-transformed control plant lines and in up to twelve independent transformed lines for each sense construct and each anti-sense construct described in Example 5 was determined as described in Example 3. The column labeled "TGA" in Table 3 shows the thioglycolic acid extractable lignins for all plant lines  
30 assayed, expressed as the average percentage of TGA extractable lignins in transformed plants versus control plants. The range of variation is shown in parentheses.

TABLE 4

	transgene	orientation	no. of lines	Northern	TGA
5	control	na	3	blank	100 (92-104)
	C3H	sense	5	3.7E+4	74 (67-85)
	F5H	sense	10	5.8E+4	70 (63-79)
	F5H	anti-sense	9	5.8E+4	73 (35-93)
10	CCR	sense	1	na	74
	CCR	anti-sense	2	na	74 (62-86)
	transgene	orientation	no. of lines	Northern	TGA
	PAL	sense	5	1.9E+5	77 (71-86)
	PAL	anti-sense	4	1.5E+4	62 (37-77)
15	C4H	anti-sense	10	5.8E+4	86 (52-113)
	PNL	anti-sense	6	1.2E+4	88 (70-114)
	LAC	sense	5	1.7E+5	na
	LAC	anti-sense	12	1.7E+5	88 (73-114)

20 Transformed plant lines containing the sense and the anti-sense lignin biosynthetic gene constructs all exhibited significantly decreased levels of lignin, relative to the empty vector-transformed control plant lines. The most dramatic effects on lignin concentration were seen in the F5H anti-sense plants with as little as 35% of the amount of lignin in control plants, and in the PAL anti-sense plants with as little as 37% of the amount of

25 lignin in control plants. These data clearly indicate that the concentration of a polynucleotide, such as lignin, as measured by the TGA assay, can be directly manipulated by conventional anti-sense methodology and also by sense over-expression using the inventive lignin biosynthetic genes, starting the whole process from an isolated cDNA obtained as in Example 1.

30

### Example 6

#### Modulation of Lignin Enzyme Activity in Transformed Plants

35

The activities and substrate specificities of selected lignin biosynthetic enzymes were assayed in crude extracts from transformed tobacco plants containing sense and anti-sense constructs for PAL (SEQ ID NO: 60), PNL (SEQ ID NO: 64) and LAC (SEQ ID NO: 65) from *Pinus radiata*, and CGT (SEQ ID NO: 59) from *Eucalyptus grandis*.

40

Enzyme assays were performed using published methods for PAL (Southerton SG and Deverall BJ, *Plant Path.* 39:223-230, 1990); CGT (Vellekoop *et al.*, *FEBS Lett.*

330:36-40, 1993); PNL (Espin *et al.*, *Phytochemistry* 44:17-22, 1997); and LAC (Bao *et al.*, *Science* 260:672-674, 1993). The data shown in the column labelled "Enzyme" in Table 5 shows the average enzyme activity from replicate measures for all plant lines assayed, expressed as a percent of enzyme activity in empty vector-transformed control plants. The range of variation is shown in parentheses.

TABLE 5

	Transgene	orientation	no. of lines	enzyme
10	control	na	3	100
	PAL	sense	5	87 (60-124)
	PAL	anti-sense	3	53 (38-80)
	CGT	anti-sense	1	89
15	PNL	anti-sense	6	144 (41-279)
	LAC	sense	5	78 (16-240)
	LAC	anti-sense	11	64 (14-106)

20 All of the transformed plant lines, except the PNL anti-sense transformed plant lines, showed average enzyme activities that were significantly lower than the activities observed in empty vector-transformed control plants. The most dramatic effects on lignin enzyme activities were seen in the PAL anti-sense transformed plant lines, in which all of the lines showed reduced PAL activity, and in the LAC anti-sense transformed plant lines, which showed as little as 14% of the LAC activity in empty vector-transformed control plant lines. These results demonstrate that enzyme activity can be modulated by transforming plants with polynucleotides encoding an enzyme of interest, starting the whole process from polynucleotides encoding enzymes of interest isolated from cDNA as described in Example 1.

### Example 7

#### Functional Identification of Lignin Biosynthetic Genes

35 Sense constructs containing DNA sequences including the coding regions for PAL (SEQ ID NO: 61), OMT (SEQ ID NO: 54), 4CL (SEQ ID NOS: 55 and 66) and POX (SEQ ID NO: 67) from *Pinus radiata*, and OMT (SEQ ID NOS: 68 and 69), CCR (SEQ ID NOS: 70 - 72), CGT (SEQ ID NOS: 59 and 73) and POX (SEQ ID NOS: 74 and 75)

from *Eucalyptus grandis* were inserted into the commercially available protein expression vector, pProEX-HT (Gibco BRL). The resultant constructs were transformed into *E. coli* XL1-Blue (Stratagene), which were then induced to produce recombinant protein by the addition of IPTG. Purified proteins were produced for the *Pinus* OMT and 4CL constructs  
 5 and the *Eucalyptus* OMT and POX constructs using Ni column chromatography (Janknecht *et al.*, *Proc. Natl. Acad. Sci. USA* 88:8972-8976, 1991). Enzyme assays for each of the purified proteins conclusively demonstrated the expected substrate specificity and enzymatic activity for the genes tested.

The data for two representative enzyme assay experiments, demonstrating the  
 10 verification of the enzymatic activity of a *Pinus radiata* 4CL gene (SEQ ID NO: 55) and a *Pinus radiata* OMT gene (SEQ ID NO: 54), are shown below in Table 6. For the 4CL enzyme, one unit equals the quantity of protein required to convert the substrate into product at the rate of 0.1 absorbance units per minute. For the OMT enzyme, one unit equals the quantity of protein required to convert 1 pmole of substrate to product per  
 15 minute.

TABLE 6

20	transgene	purification step	total ml extract	total mg protein	total units activity	% yield activity	fold purification
	4CL	crude	10 ml	51 mg	4200	100	1
		Ni column	4 ml	0.84 mg	3680	88	53
25	OMT	crude	10 ml	74 mg	4600	100	1
		Ni column	4 ml	1.2 mg	4487	98	60

The data shown in Table 6 demonstrate that both the purified 4CL enzyme and the  
 30 purified OMT enzyme show high activity in enzyme assays, confirming the identification of the 4CL and OMT genes. Crude protein preparations from *E. coli* transformed with empty vector show no activity in either the 4CL or the OMT enzyme assay. This demonstrates that the function of an isolated novel cDNA with only a putative function can be confirmed, starting the whole process from an isolated cDNA obtained as in  
 35 Example 1.

### Example 8

#### Demonstration of the Presence / Absence of Unique Sequence Identifiers in Plants

5 Transgenic tobacco plants were created using unique identifier sequences which are not found in tobacco. The unique identifier sequences inserted were isolated from *Pinus radiata*, SEQ ID NO: 76, and *Eucalyptus grandis*, SEQ ID NO: 77. The unique identifier sequences were inserted into *Agrobacterium tumefaciens* LBA4301 (provided as a gift by Dr. C. Kado, University of California, Davis, CA) by direct transformation using  
10 published methods (An *et al.*, "Binary vectors," in Gelvin SB and Schilperoort RA, eds., *Plant Molecular Biology Manual*, Kluwer Academic Publishers: Dordrecht, 1988). The presence and integrity of the unique identifier sequences in the *Agrobacterium* transgenic constructs were verified by restriction digestion and DNA sequencing.

Tobacco (*Nicotiana tabacum* cv. Samsun) leaf sections were transformed using the  
15 method of Horsch *et al.*, *Science* 227:1229-1231, 1985. Three independent transformed plant lines were established for each unique sequence identifier used. Two empty-vector control plant lines were established using an empty gene transfer vector that lacked a unique sequence identifier.

The uniqueness of the sequence identifiers was assayed using Southern blot  
20 analyses to test for the presence of the sequence identifier in the genome of the plants. If the sequence identifier is unique and therefore useful as a tag, then the sequence identifier should be clearly absent in plants which have not been tagged and it should be clearly present in plants which have been tagged. In the present example, the unique identifiers would be expected to be absent in the empty-vector transformed control plants. The  
25 unique identifier would be expected to be present in the transgenic plants transformed with the unique sequence identifiers.

Genomic DNA was prepared from empty-vector transformed control plants and plants transformed with unique sequence identifiers using the cetyltrimethyl-ammonium bromide (CTAB) extraction method of Murray MG and Thompson WF, *Nucleic Acids*  
30 *Res.* 8:4321-4325, 1980. The DNA samples were digested with the restriction enzyme *EcoRI* in the case of the plants transformed with the *Pinus* unique sequence identifier (SEQ ID NO: 76) and the restriction enzyme *XbaI* in the case of the plants transformed with the *Eucalyptus* unique sequence identifier (SEQ ID NO: 77). The DNA fragments

produced in the restriction digests were resolved on a 1% agarose gel; the left panel of Fig. 2 and the right panel of Fig. 2 show the DNA fragment patterns of the DNA samples from the *Pinus* and *Eucalyptus* experiments, respectively.

After the agarose gel electrophoresis step, the DNA samples were transferred to  
5 Hybond-N+ nylon membranes (Amersham Life Science, Little Chalfont, Buckinghamshire, England) using methods established by Southern, *J. Mol. Biol.* 98:503-517, 1975. The nylon membranes were probed with radioactively-labeled probes for the unique sequence identifiers identified above and washed at high stringency (final wash: 0.5 X salt sodium citrate buffer (SSC) plus 0.1% sodium dodecyl sulfate (SDS), 15  
10 minutes at 65°C). The hybridization of the probes to complementary sequences in the genomic DNA samples was detected using auto-radiography.

The results are shown in Figs. 3 and 4.

Fig. 3 (corresponding to the left panel of Fig. 2) shows the hybridization pattern detected in the Southern blot analysis using a probe derived from the *Pinus* sequence  
15 identifier (SEQ ID NO: 76). Lanes A-B contain DNA samples from empty-vector transformed control plants and Lanes C-E contain DNA from plants transformed with SEQ ID NO: 76. There is no hybridization in Lanes A-B indicating that SEQ ID NO: 76 is not present in empty-vector transformed tobacco plants; that is, SEQ ID NO: 76 is a unique tag suitable for unambiguous marking of tobacco plants. There is strong  
20 hybridization in Lanes C-E, indicating that the plants which received SEQ ID NO: 76 via transformation have been clearly and unambiguously tagged with the unique sequence contained in SEQ ID NO: 76.

Fig. 4 (corresponding to the right panel of Fig. 2) shows the hybridization pattern detected in the Southern blot analysis using a probe derived from the *Eucalyptus* sequence  
25 identifier (SEQ ID NO: 77). Lanes A-B contain DNA samples from empty-vector transformed control plants and Lanes C-E contain DNA from plants transformed with SEQ ID NO: 77. There is no hybridization in Lanes A-B indicating that SEQ ID NO: 77 is not present in empty-vector transformed tobacco plants; that is, SEQ ID NO: 77 is a unique tag suitable for unambiguous marking of tobacco plants. There is strong  
30 hybridization in Lanes C-E indicating that the plants which received SEQ ID NO: 77 via transformation have been clearly and unambiguously tagged with the unique sequence contained in SEQ ID NO: 77.

The data clearly demonstrates the utility of the sequences disclosed in this specification for the purposes of unambiguously tagging transgenic materials. A unique sequence was selected from a large number of potential tags and shown to be absent in the genome of the organism to be tagged. The tag was inserted into the genome of the  
5 \_ organism to be tagged and a well-established DNA detection method was used to clearly detect the unique sequence identifier used as the tag.

Because of the sequence-specific detection methods used in the example, a user of the invention disclosed in this specification has both a high likelihood of finding a sequence identifier, among the list which has been disclosed, which will be useful for  
10 tagging any given organism and an unequivocal method for demonstrating that a tagged organism could only have acquired a given tag through the deliberate addition of the unique sequence to the genome of the organism to be tagged. If the user of this invention maintains the precise sequence of the tag used in a given organism as a secret, then any disputes as to the origin and history of the organism can be unambiguously resolved using  
15 the tag detection techniques demonstrated in the present example.

SEQ ID NOS: 1-304 are set out in the attached Sequence Listing. The codes for nucleotide sequences used in the attached Sequence Listing, including the symbol "n," conform to WIPO Standard ST.25 (1998), Appendix 2, Table 1.

All references cited herein, including patent references and non-patent  
20 publications, are hereby incorporated by reference in their entirety. While in the foregoing specification, this invention has been described in relation to certain preferred embodiments, and many details have been set forth for purposes of illustration, it will be apparent to those skilled in the art that the invention is susceptible to additional embodiments and that certain of the details described herein may be varied considerably  
25 without departing from the basic principles of the invention.



Claims:

1. An isolated polynucleotide comprising a polynucleotide sequence selected from the group consisting of: (1) the sequences recited in SEQ ID NOS: 1-53 and 78-164; (2) complements of the sequences recited in SEQ ID NOS: 1-53 and 78-164; (3) reverse complements of the sequences recited in SEQ ID NOS: 1-53 and 78-164; (4) reverse sequences of the sequences recited in SEQ ID NOS: 1-53 and 78-164; (5) sequences comprising a polynucleotide sequence having at least 40% identity to a compare sequence selected from the polynucleotide sequences recited in SEQ ID NOS: 1, 2, 4-6, 8-12, 19, 21-23, 28-33, 35, 37-42, 44, 46-52, 78-80, 82, 83, 86, 89-92, 96-100, 104-113, 115, 117, 120, 122-130, 132-136, 138-158, 160, 163 and 164, the percentage identity determined by aligning the sequence and the compare sequences using the BLASTN algorithm version 2.04 set at the parameter values described herein, identifying the number of identical nucleic acids over aligned portions of the sequence and the compare sequences, dividing the number of identical nucleic acids by the total number of nucleic acids of the compare sequence, and multiplying by 100 to determine the percentage identity; (6) sequences comprising a polynucleotide sequence having at least 60% identity to a compare sequence selected from the polynucleotide sequences recited in SEQ ID NOS: 3, 7, 14, 18, 20, 25, 34, 36, 53, 84, 85, 87, 88, 101, 114, 116, 118, 119, 131, 137, 159, 161 and 162, the percentage identity determined as described in (5) above; (7) sequences comprising a polynucleotide sequence having at least 75% identity to a compare sequence selected from the polynucleotide sequences recited in SEQ ID NOS: 16, 17, 26, 43, 45, 93, 94 and 121, the percentage identity determined as described in (5) above; (8) sequences comprising a polynucleotide sequence having at least 90% identity to a compare sequence selected from the nucleotide sequences recited in SEQ ID NOS: 13, 24, 95, 102 and 103, the percentage identity determined as described in (5) above; (9) sequences comprising a polynucleotide sequence that hybridizes to a polynucleotide comprising a sequence recited in (1) – (8) above under stringent hybridization conditions; (10) sequences comprising a polynucleotide sequence that is a 100-mer of a sequence recited in (1) – (8) above; (11) sequences comprising a polynucleotide sequence that is a 40-mer of a sequence recited in (1) – (8) above; and (12) sequences

- comprising a polynucleotide sequence that is a 20-mer of a sequence recited in (1) – (8) above; and (13) sequences comprising a polynucleotide sequence differing from a sequence recited in (1) – (12), above, only by one or more conservative substitutions.
- 5    2.    An isolated oligonucleotide probe or primer comprising at least 10 contiguous residues complementary to 10 contiguous residues of a nucleotide sequence recited in Claim 1.
3.    A genetic construct comprising a polynucleotide described in claim 1.
4.    A transgenic cell comprising a genetic construct according to claim 3.
- 10   5.    A transgenic cell according to claim 4, wherein the cell is selected from one of the following: a bacterial cell; an insect cell; a yeast cell; a mammalian cell; and a plant cell.
6.    A genetic construct comprising, in the 5'-3' direction:
- 15       (a)    a gene promoter sequence;
- (b)    a polynucleotide sequence comprising at least one of the following: (1) a polynucleotide comprising a nucleotide sequence of claim 1 coding for at least a functional portion of an enzyme having activity in an isoprenoid biosynthetic pathway; and (2) a polynucleotide comprising nucleotide sequence of claim 1 that includes a non-coding region of a polynucleotide encoding an enzyme having activity in an isoprenoid biosynthetic pathway;
- 20           and
- (c)    a gene termination sequence.
7.    The construct of claim 6 wherein the polynucleotide is in a sense orientation.
8.    The construct of claim 6 wherein the polynucleotide is in an antisense orientation.
- 25   9.    The construct of claim 6 wherein the gene promoter sequence and gene termination sequences are functional in a plant host.
10.   A transgenic cell comprising a construct of claim 6.
11.   The transgenic cell of claim 10 wherein the polynucleotide is in a sense orientation.
- 30   12.   The transgenic plant cell of claim 10 wherein the polynucleotide is in an antisense orientation.

13. A transgenic cell according to claim 10, wherein the cell is selected from one of the following: a bacterial cell; an insect cell; a yeast cell; a mammalian cell; and a plant cell.
14. A plant comprising a transgenic cell according to claim 9, or fruit or seeds or progeny thereof.
15. The plant of claim 14 wherein the plant is a woody plant.
16. The plant of claim 15 wherein the plant is selected from the group consisting of eucalyptus and pine species.
17. A method for modulating one or more of the content, the composition, and the metabolism of an enzyme involved in an isoprenoid biosynthetic pathway in an organism, comprising stably incorporating into the genome of the organism a construct of claim 3.
18. A method according to claim 17, wherein the organism is a plant.
19. A method for modulating one or more of the content, the composition, and the metabolism of an isoprenoid compound in an organism comprising stably incorporating into the genome of the organism a construct of claim 6.
20. A method according to claim 19, wherein the organism is a plant.
21. A method for producing an organism having one or more of altered isoprenoid content, altered isoprenoid composition and altered isoprenoid metabolism, comprising:
- (a) transforming a host cell with a construct of claim 3 to provide a transgenic host cell; and
  - (b) cultivating the transgenic host cell under conditions conducive to growth and regeneration.
22. A method according to claim 21, wherein the organism is a plant and the host cell is a plant cell.
23. An isolated polypeptide encoded by a polynucleotide of claim 1.
24. A polypeptide of claim 23 having enzymatic activity in an isoprenoid biosynthetic pathway in a plant.
25. An isolated polypeptide comprising an amino acid sequence expressed from a polynucleotide that hybridizes to a nucleotide sequence set forth as SEQ ID NOS: 1-53 and 78-164 under stringent hybridization conditions.

26. An isolated polypeptide comprising a polypeptide sequence selected from the group consisting of: (1) the sequences set forth in SEQ ID NOS: 165-286 and 288-304; (2) sequences comprising a polypeptide sequence having at least 50% identity to a compare sequence selected from the polypeptide sequences recited in  
5 SEQ ID NOS: 194-200, 202, 216, 223, 230, 235, 239, 240, 243, 250, 255, 259, 260, 263, 270, 272, 274, 278, 291, 292, 293, 296, 303 and 304; (3) sequences comprising a polypeptide sequence having at least 75% identity to a compare sequence selected from the polypeptide sequences recited in SEQ ID NOS: 166, 168-177, 179, 183-188, 192, 203-205, 207, 209-213, 218, 219, 221, 224, 225, 227-  
10 229, 231, 232, 234, 237, 242, 244, 245, 251, 253, 262, 267, 268, 269, 273, 276, 277, 279, 281, 282, 284, 286, 289, 290, 294, 295, and 297-302; (4) sequences comprising a polypeptide sequence having at least 90% identity to a compare sequence selected from the polypeptide sequences recited in SEQ ID NOS: 165, 167, 178, 182, 189-191, 193, 201, 206, 208, 214, 215, 217, 220, 222, 226, 233,  
15 238, 241, 246-250, 254, 256-258, 261, 264, 265, 266, 275, 280, 283, 285 and 288; (5) sequences comprising a polypeptide sequence having at least 95% identity to a compare sequence selected from the polypeptide sequences recited in SEQ ID NOS: 180, 181 and 271; (6) sequences comprising a polypeptide sequence that is a 100-mer of a sequence recited in (1) – (5) above having at least 100 residues; (7)  
20 sequences comprising a polypeptide sequence that is a 40-mer of a sequence recited in (1) – (5) above having at least 40 residues; and (8) sequences comprising a polypeptide sequence that is a 20-mer of a sequence recited in (1) – (5) above.
27. A method for modulating one or more of the content, the composition and the metabolism of an isoprenoid compound in an organism, comprising administering  
25 an isolated polypeptide of claim 26 to the organism.
28. A method according to claim 27, wherein the organism is a plant, and administration of the isolated polypeptide is topical.
29. A method according to claim 27, wherein the organism is a mammal, and administration of the isolated polypeptide is systemic.

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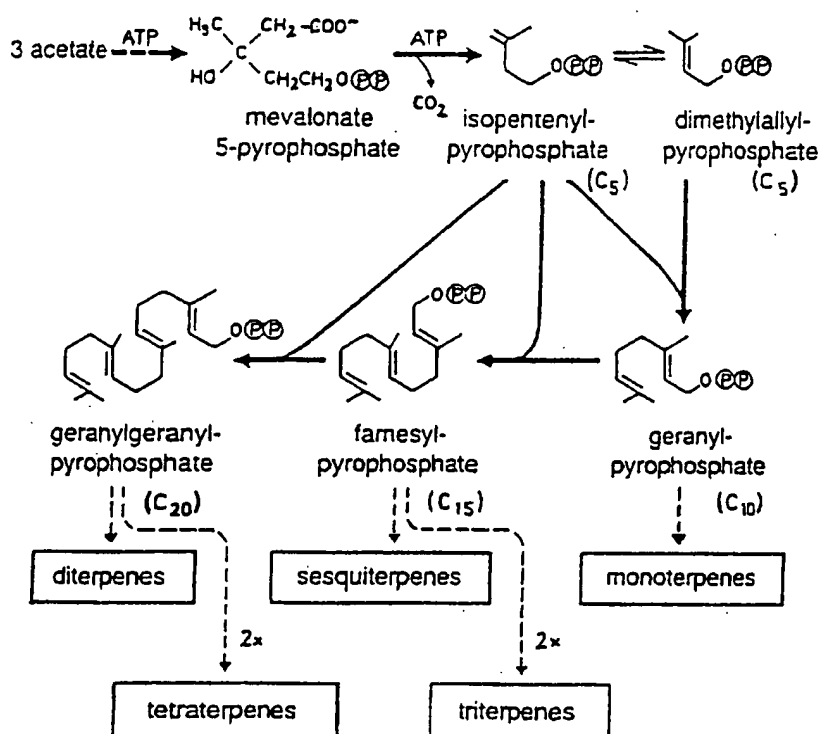


FIGURE 1

2/4

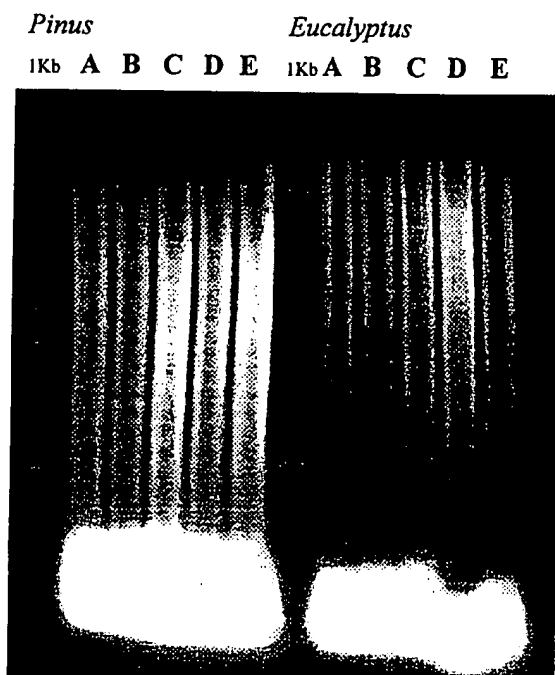


FIGURE 2

3/4

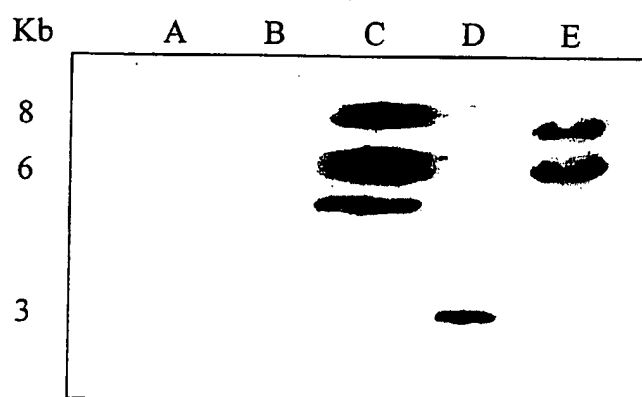


FIGURE 3

4/4

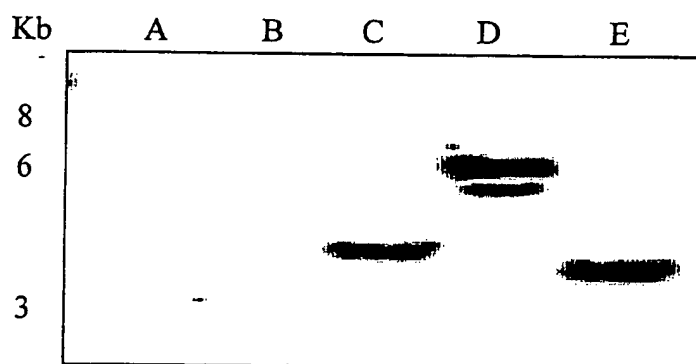


FIGURE 4



## SEQUENCE LISTING

&lt;110&gt; Havukkala, Iilka

<120> Materials and Methods for the Modification  
of Isoprenoid Content, Compostition and Metabolism

&lt;130&gt; 11000.1019c1PCT

&lt;160&gt; 304

&lt;170&gt; FastSEQ for Windows Version 3.0

&lt;210&gt; 1

&lt;211&gt; 414

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 1

gatattaaga aaattgtgga gctgatgtct gatctgcatt ttatttataa cacgcacaga	60
ttcgcttatt tgtattcgaa attcaatagc tctatttaca tgtataagtt cagtttggac	120
actgacctta atattgtcaa gaaaatgagt gggttcgacg tagaagggtg atgtcatgct	180
gacgagttat tctacttttt ctccacaaac atgacgaaag actactacga atcggaggac	240
aaaatcaaag aatatgtgtg gaaggtgacg aaactgtgga caaacttcgc taaaaccagt	300
aacccaactc cagacacgtc actaggcgtg tcttggccga ggtacaccat ggctaacaag	360
gaatacctgg acatcaacac gcagctaaca acgggacgct actcggagcg ggaa	414

&lt;210&gt; 2

&lt;211&gt; 1834

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 2

attgcctgct tctgcttcaa ttgaagcttt tctgctcgcc aatcaatatg gcaattgcaa	60
gcagggcagg agttgcaccc atccttcaag tagactgtca ttccacccat ttcaattcaa	120
tgactgaatt gggttctcgg aatagtatga tgtttcagtc tgcaattccc tgctcatttc	180
gacagattag ggcgaccact aaaagaaagc gttgtgtctt gttggctaag ctaagtaatt	240
ccgatggaga aaatgggaaa aatgtgaagg cagctgtgga gattgcttca aagagtggat	300
tcccagctga gaaacctcct acaccgttgc ttgacaccgt taattatcca gtacacttaa	360
agaatctctc tatacaggat cttagagcaac tggccacaga aattagagca gaacttgtgt	420
tcggtgtggc aaaaactgga ggccacctgg gaggtagcct ggggtgtggta gatctaacag	480
tggtcttcca ccatgtcttt gattctccag aggataggat tatatgggat gttggtcacc	540
agtcatatcc acacaagatt ttgacaggga gaagatccaa aatgcataca atcagacaga	600
cctctggttt ggccggattt cccaaacggg acgaaaagcaa atacgatgct ttggcgctg	660
gacatagttc taccagtatc tccgctggac tcggtatggc agttgggaga gattttattga	720
agaaaaataa ccatgtgggtg gccgtgattg gggatggagc catgacagca ggacaagctt	780
acgaggctat gaataattcg ggatatctgg aatccaatct tatcataatt cttaatgata	840
ataagcaagt ttctctgcca actgccacac tagacggagc tgcgcgcgcc gtaggtgcac	900
ttaccagggc tctcaciaag ctccagtcca gcaaaaaact tcgcaaaact agagaagctg	960
ccaaggtct caccaagcag ataggtgggc caactcatga agtagcatcg aaagttgata	1020
agtatgcaag gggctcttct agcccagcaa gttcctcgtt gtttgacgag ctaggtcttt	1080
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gttacacctc agcggaggag gctgcagaca aattacatgg tgtggtaaag ttcgatccag	1260
ttacgggcaa gcaattcaa tcaaaaagtt ccgttctgag ctatacacag tactttgcag	1320
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<400> 4						
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 <213> Pinus radiata

<400> 5						
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 <212> DNA  
 <213> Eucalyptus grandis

<400> 6						
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&lt;210&gt; 7

&lt;211&gt; 699

&lt;212&gt; DNA

<213> *Eucalyptus grandis*

&lt;400&gt; 7

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&lt;210&gt; 8

&lt;211&gt; 373

&lt;212&gt; DNA

<213> *Pinus radiata*

&lt;400&gt; 8

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&lt;210&gt; 9

&lt;211&gt; 373

&lt;212&gt; DNA

<213> *Pinus radiata*

&lt;400&gt; 9

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&lt;210&gt; 10

&lt;211&gt; 825

&lt;212&gt; DNA

<213> *Eucalyptus grandis*

&lt;400&gt; 10

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&lt;210&gt; 11

&lt;211&gt; 394

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 11

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&lt;210&gt; 12

&lt;211&gt; 245

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 12

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&lt;210&gt; 13

&lt;211&gt; 375

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 13

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&lt;210&gt; 14

&lt;211&gt; 824

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 14

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&lt;211&gt; 1271

&lt;212&gt; DNA

<213> *Eucalyptus grandis*

&lt;400&gt; 15

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&lt;210&gt; 16

&lt;211&gt; 372

&lt;212&gt; DNA

<213> *Pinus radiata*

&lt;400&gt; 16

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&lt;210&gt; 17

&lt;211&gt; 520

&lt;212&gt; DNA

<213> *Pinus radiata*

&lt;400&gt; 17

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&lt;210&gt; 18

&lt;211&gt; 435

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 18

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&lt;210&gt; 19

&lt;211&gt; 320

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 19

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&lt;210&gt; 20

&lt;211&gt; 626

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 20

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&lt;210&gt; 21

&lt;211&gt; 490

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 21

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tctttgacaa						490

&lt;210&gt; 22

&lt;211&gt; 396

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 22

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&lt;210&gt; 23

&lt;211&gt; 396

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 23

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&lt;210&gt; 24

&lt;211&gt; 700

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 24

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&lt;210&gt; 25

&lt;211&gt; 1513

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 25

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&lt;210&gt; 26

&lt;211&gt; 295

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 26

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gagatgggtg aaagaatcgg gttccctga gctaaccctc gctcgacatc gttacgtgga      180
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&lt;210&gt; 27

&lt;211&gt; 191

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 27

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gctgtaataa caattgatat gctatgcata tatttgaata aacaaatact cgttggccat      180
ctattttatt t                                     191

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&lt;210&gt; 28

&lt;211&gt; 373

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 28

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agcgatactt	gga					373

&lt;210&gt; 29

&lt;211&gt; 1411

&lt;212&gt; DNA

<213> *Eucalyptus grandis*

&lt;400&gt; 29

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ttgtggccaa	gctaattctc	gcgttgctga	ttccgagatc	cggaaagcgc	ctccctcccc	240
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&lt;210&gt; 30

&lt;211&gt; 689

&lt;212&gt; DNA

<213> *Eucalyptus grandis*

&lt;400&gt; 30

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&lt;210&gt; 31

&lt;211&gt; 393

&lt;212&gt; DNA

<213> *Pinus radiata*

&lt;400&gt; 31

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&lt;210&gt; 32

&lt;211&gt; 519

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 32

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&lt;210&gt; 33

&lt;211&gt; 302

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 33

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aa						302

&lt;210&gt; 34

&lt;211&gt; 508

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 34

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&lt;210&gt; 35

&lt;211&gt; 353

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 35

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&lt;210&gt; 36

&lt;211&gt; 82

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 36

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&lt;210&gt; 37

&lt;211&gt; 474

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 37

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&lt;210&gt; 38

&lt;211&gt; 340

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 38

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&lt;210&gt; 39

&lt;211&gt; 487

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 39

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&lt;210&gt; 40

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<210> 41  
 <211> 512  
 <212> DNA  
 <213> Eucalyptus grandis

<400> 41  
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 tttggtggat gaagaggatc atgtcattgg gcatgactca aaatacaatt gtcacttgat 360  
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 aaaatatgaa ttgcttcttc agcaa 445

<210> 43  
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 <212> DNA  
 <213> Eucalyptus grandis

<400> 43  
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 cggcgccctc tgttcgagga cgaatgcac ttggtggatg agaacgacaa tgcctgtcgt 180  
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 gccacaaagg taacattccc ccttgtgtgg acaaacacct gctgcagcca tccattgtac 360  
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<210> 44  
 <211> 834  
 <212> DNA  
 <213> Pinus radiata

<400> 44

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<210> 45  
 <211> 389  
 <212> DNA  
 <213> Pinus radiata

<400> 45

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cctcaagaag	aagatcggcc	ttgtctatca	gctcaacatt	tcaccaaga	aaattggaat	300
agctgaggag	gtgttcgttg	tggacctcaa	gaatggcaaa	gtcactaaag	gaccatatga	360
aggaaagcca	gatgcaacat	tttcctttg				389

<210> 46  
 <211> 469  
 <212> DNA  
 <213> Pinus radiata

<400> 46

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tttccaaagg	gaaaaggaga	gatcaatgag	gaaggagtag	cctattacaa	taacctcatc	360
aatgaactcc	tccagaatgg	aatccaagcg	tctgtcactt	tgtttctactg	ggatactccc	420
cagtctcttg	aggatgaata	tggcggattt	ctgaggccaa	ccattgtga		469

<210> 47  
 <211> 349  
 <212> DNA  
 <213> Pinus radiata

<400> 47

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349

<210> 48  
 <211> 385  
 <212> DNA  
 <213> Pinus radiata

<400> 48  
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 ctgaagtga taaggaagga atagc 385

<210> 49  
 <211> 417  
 <212> DNA  
 <213> Pinus radiata

<220>  
 <221> unsure  
 <222> (209) ... (212)

<400> 49  
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 aagatatggg agtggatgtc tacagattct caatctcatg gtctcgaatg tttccaaaag 360  
 gaaaagggga gatcaatgag gaaggagtag cctattacaa taacctcata aatgaac 417

<210> 50  
 <211> 264  
 <212> DNA  
 <213> Pinus radiata

<400> 50  
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 gatcaagtac tgggggaaaa gagatgaaaa gctgatcctt cccatcaatg acagcatcag 180  
 ctttactttg gatccagacc atctgtcagc cacaaccact gtagcagtta gccatcatt 240  
 cacatctgat agaattgtggc tcaa 264

<210> 51  
 <211> 417  
 <212> DNA  
 <213> Eucalyptus grandis

<400> 51  
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 tcaaggttga gacctgatca gagacactta gaaaatgggg aatatgagtt ggcaaatgcc 180  
 gagaagttaa gactggaaca catacagaga caggcaagaa agttacagga gggaggttgg 240  
 caaccgagat gggttgggaa ggatgatgat ggatgttacc gctacatggg tgggtattgg 300  
 gaagctcgag aagcatacga actgggatgg aatccctgac atattcgggc aaaaatgttg 360  
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<210> 52

<211> 305  
 <212> DNA  
 <213> Pinus radiata

<400> 52  
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 gcagaaaaatg aacattttac atacagtctg tcctcaaaag taaaaaccaa gtttcttggc 180  
 aactctgtgg atattttacc acttggaagg acacgtgtgg tgctaaagaa atccggagac 240  
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 attga 305

<210> 53  
 <211> 474  
 <212> DNA  
 <213> Eucalyptus grandis

<400> 53  
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 atggagcaga tgaagcagca cctctccacc gagcgcggca aggcgggtcac caagaagatc 180  
 ggcctcgtct accagatcaa catcgccccc aagaaaattg ggttcgacga ggtggtctac 240  
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 caaattgctt ttatgagggg agcaatgaag attaaggga gcttgagtgc agcgcagaaa 420  
 ttcactcctg acatattccc aaagccatcg aagatgtgag cattttgaaa agg 474

<210> 54  
 <211> 562  
 <212> DNA  
 <213> Pinus radiata

<400> 54  
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 cgtgtaccct cgtgagcccg agccaatgaa ggagctccgc gaagtgactg ccaagcatcc 360  
 ctggaacctc atgactactt ctgccgatga ggggtcaattt ctgggcctcc tgctgaagct 420  
 cattaacgcc aagaacacca tggagattgg ggtgtacact ggttactcgc ttctcagcac 480  
 agcccttgca ttgcccgatg atggaaagat tctagccatg gacatcaaca gagagaacta 540  
 tgatatcgga ttgcctataa tt 562

<210> 55  
 <211> 1961  
 <212> DNA  
 <213> Pinus radiata

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 aagccgacga aaccatgc cggccgtga caatccacc ggacgatgtc gtggcggttc 660

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&lt;210&gt; 56

&lt;211&gt; 414

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 56

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&lt;210&gt; 57

&lt;211&gt; 469

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 57

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&lt;210&gt; 58

&lt;211&gt; 760

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 58

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&lt;210&gt; 59

&lt;211&gt; 468

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 59

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&lt;210&gt; 60

&lt;211&gt; 684

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 60

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&lt;210&gt; 61

&lt;211&gt; 479

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 61

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&lt;210&gt; 62

&lt;211&gt; 1785

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 62

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&lt;210&gt; 63

&lt;211&gt; 475

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 63

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&lt;210&gt; 64

&lt;211&gt; 957

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 64

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&lt;210&gt; 65

&lt;211&gt; 471

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 65

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&lt;210&gt; 66

&lt;211&gt; 1010

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 66

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&lt;210&gt; 67

&lt;211&gt; 1410

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 67

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gacaactaca	tatattcttt	aaaaaaaaaa				1410

&lt;210&gt; 68

&lt;211&gt; 607

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 68

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gccggtg						607

&lt;210&gt; 69

&lt;211&gt; 421

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 69

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c						421

&lt;210&gt; 70

&lt;211&gt; 508

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 70

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&lt;210&gt; 71

&lt;211&gt; 495

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 71

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&lt;210&gt; 72

&lt;211&gt; 472

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 72

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&lt;210&gt; 73

&lt;211&gt; 380

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 73

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&lt;210&gt; 74

&lt;211&gt; 515

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 74

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&lt;210&gt; 75

&lt;211&gt; 487

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 75

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gagagg						487

&lt;210&gt; 76

&lt;211&gt; 1474

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 76

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&lt;210&gt; 77

&lt;211&gt; 414

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 77

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&lt;210&gt; 78

&lt;211&gt; 273

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 78

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&lt;210&gt; 79

&lt;211&gt; 121

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 79

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&lt;210&gt; 80

&lt;211&gt; 505

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 80

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&lt;210&gt; 81

&lt;211&gt; 270

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 81

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 <212> DNA  
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<400> 82  
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 <213> *Eucalyptus grandis*

<400> 83  
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<210> 84  
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 <212> DNA  
 <213> *Pinus radiata*

<400> 84  
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<210> 85  
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 <212> DNA  
 <213> *Pinus radiata*

<400> 85  
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<210> 86  
 <211> 247



&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 86

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&lt;210&gt; 87

&lt;211&gt; 426

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 87

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gttcct						426

&lt;210&gt; 88

&lt;211&gt; 488

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 88

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&lt;210&gt; 89

&lt;211&gt; 223

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 89

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&lt;210&gt; 90

&lt;211&gt; 318

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 90

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 <212> DNA  
 <213> Eucalyptus grandis

<400> 91

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 <212> DNA  
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<400> 92

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 <213> Eucalyptus grandis

<400> 93

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<210> 94  
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<400> 94						
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<400> 96						
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<400> 97						
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&lt;210&gt; 98

&lt;211&gt; 668

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 98

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&lt;210&gt; 99

&lt;211&gt; 430

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 99

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gtgttgaacg						430

&lt;210&gt; 100

&lt;211&gt; 478

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 100

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 <213> Pinus radiata

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 <212> DNA  
 <213> Pinus radiata

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 <212> DNA  
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&lt;400&gt; 105

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&lt;210&gt; 106

&lt;211&gt; 265

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 106

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catctccgcc	ggctcggta	tgggc				265

&lt;210&gt; 107

&lt;211&gt; 295

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 107

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&lt;210&gt; 108

&lt;211&gt; 456

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 108

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&lt;210&gt; 109

&lt;211&gt; 640

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 109

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ccgcgggtta	tcacccagc	gtatgggggg	actacttct	taaatatgat	tctccctcca	180
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&lt;210&gt; 110

&lt;211&gt; 396

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 110

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caaatccaac	gcttgggaat	tgagtaccat	tttgaacgtg	aaatagatga	gcaattagaa	360
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&lt;210&gt; 111

&lt;211&gt; 348

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 111

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&lt;210&gt; 112

&lt;211&gt; 508

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 112

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&lt;210&gt; 113

&lt;211&gt; 398

&lt;212&gt; DNA

<213> *Eucalyptus grandis*

&lt;400&gt; 113

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&lt;210&gt; 114

&lt;211&gt; 432

&lt;212&gt; DNA

<213> *Pinus radiata*

&lt;400&gt; 114

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&lt;210&gt; 115

&lt;211&gt; 363

&lt;212&gt; DNA

<213> *Pinus radiata*

&lt;400&gt; 115

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cat						363

&lt;210&gt; 116

&lt;211&gt; 779

&lt;212&gt; DNA

<213> *Pinus radiata*

&lt;400&gt; 116

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 <212> DNA  
 <213> Pinus radiata

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 <213> Pinus radiata

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 <212> DNA  
 <213> Eucalyptus grandis

<400> 121  
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 <212> DNA  
 <213> Eucalyptus grandis

<400> 122  
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&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 123

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&lt;210&gt; 124

&lt;211&gt; 604

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 124

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tcaa						604

&lt;210&gt; 125

&lt;211&gt; 515

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 125

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&lt;210&gt; 126

&lt;211&gt; 366

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 126

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&lt;210&gt; 127

&lt;211&gt; 458

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 127

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&lt;210&gt; 128

&lt;211&gt; 442

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 128

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&lt;210&gt; 129

&lt;211&gt; 392

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 129

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&lt;210&gt; 130

&lt;211&gt; 354

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 130

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&lt;210&gt; 131

&lt;211&gt; 442

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 131

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&lt;210&gt; 132

&lt;211&gt; 984

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 132

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&lt;210&gt; 133

&lt;211&gt; 527

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 133

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&lt;210&gt; 134

&lt;211&gt; 965

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 134

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&lt;211&gt; 503

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 135

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&lt;211&gt; 563

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 136

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&lt;210&gt; 137

&lt;211&gt; 354

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 137

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&lt;210&gt; 138

&lt;211&gt; 631

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 138

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&lt;210&gt; 139

&lt;211&gt; 362

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 139

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&lt;210&gt; 140

&lt;211&gt; 504

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 140

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tcaccagatt	gggatttgtc	atggcgacat	tgggtgaaaag	aggctggctg	tatctgatca	120
caaatttcac	tgattttcaa	ctggcttcca	taggcagttt	tcttcttcat	gagagcatct	180
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aaattcagag	caagacgaac	acagttgctg	cacaagaaaa	atgtattact	cgactgctgc	300
tatatcattt	ttgtgtcaac	ctgccagtca	tgggtggttc	ctatcctgta	ttcagattta	360
tgggcgatgac	aagcgtgcta	ccactaccat	cctggaaaag	agttgtatcc	caactggttt	420
gttatttcat	tttggaggat	tttgttttct	actggggcca	cagaatttta	cattcaaaat	480
ggctgtacaa	gcatgttcac	agtg				504

&lt;210&gt; 141

&lt;211&gt; 1293

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 141

caagtactct	ttataagctc	tgaatgattg	ctagctcaaa	taggcaacga	agaattgctt	60
gtaaactatt	cctgagaagt	gtgtctgtta	cacaattctc	aaatatcatt	gatcttcagg	120
attttgatc	acatctgaga	acccaggat	gggagaagag	ttgcagacat	ggatattaat	180
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aaagctgatc	cttcccatca	atgacagcat	cagctttact	ttggatccag	accatctgtc	300
agccacaacc	actgtagcag	ttagcccatc	attcacatct	gatagaatgt	ggctcaacgg	360
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gataggcgac	acctcaatac	taggagaaat	cgcggtggac	tcaatgaagg	atgttgaatc	1260
cttgactgct	cctccagagc	tcaagagtga	aag			1293

&lt;210&gt; 142

&lt;211&gt; 389

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 142

gtactcttta	taagctctga	atgttgctag	ctcaaatagg	cgacgaagaa	ttgcttgtaa	60
actattcctg	agaagtgggt	ctgttacaca	attctcaaac	atcattgata	ttcaggattt	120
tggtcacat	ctgagaaccc	aggtatggga	gaagagtgtc	agacatggat	attaatgggtc	180
actgcaagag	ctctacaaa	tatagcagt	atcaagtact	gggggaaaag	agatgaaaag	240
ctgatccttc	ccatcaatga	cagcatcagc	tttactttgg	atccagacca	tctgtcagcc	300
acaaccactg	tagcagttag	cccatcattc	acatctgata	gaatgtgggt	caacggcaag	360
gaggtctctc	ttggaggggg	gagatatca				389

&lt;210&gt; 143

&lt;211&gt; 693

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 143

ccgttatata	cataaatatg	ttttcttttt	aaatggcaga	gacaccctga	atggtcacat	60
gaacagagaa	gggttaaca	cgatattgcc	tgaggaccaa	gtctataaga	tagaccagga	120
tagctatgcc	tctcactctt	ctattggcga	atacatgggc	atcgtctgcc	atagtttcca	180
ggaggggtatc	gctctttgtg	gcttgctcaa	caactgtagt	ctctcgttca	ttcagtaaga	240
gctgctccgg	tgtataccc	cggagccca	aatctgctca	tcccgcactc	actgggagca	300
gaacttgctt	ctcccggaac	ccaattgtta	gaaatttgat	tggtatccgt	tctaagatgg	360
gcgcgacagt	ggaggatacg	accatggatg	ctgttcagag	gcggctcatg	ttcgaagatg	420
agtgcatttt	ggtggatgaa	gaggatcatg	tcattgggca	tgactcaaaa	tacaattgtc	480
acttgatgga	gaaaatagag	tcagagaatc	tattgcatag	agctttcagt	gtgtttctat	540
tcaatacaaaa	atatgaattg	cttcttcagc	aacgttctgc	aacaaagggtg	acattccctt	600
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agaacaattt	agggtcagaa	atgcagccca	aag			693

&lt;210&gt; 144

&lt;211&gt; 385

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 144

cgctgcagg	tcgacactag	tggatccaaa	gaagaactgg	tgtgatggca	ggaattccag	60
tcctaaggcc	atcttgcatc	tggttgcttt	cagtctacat	gctgcacatt	gtagctgcag	120
tagcttcacc	aaggctaggt	agaagcagct	tcccaagggg	tttcaaat	ggtgcagggt	180
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atcaatacca	ccgttataag	gaagatgtgc	agcttctcaa	atacatggga	atggacgtct	360
atcgtttctc	tatctcctgg	tcacg				385

&lt;210&gt; 145

&lt;211&gt; 385



&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 145

aaggccccag	catttgggat	acattctccc	acactccagg	taaaatcgct	gatggaaaga	60
atggggacgt	tgacgtagac	caataccacc	gttataagga	agatgtgcag	cttctcaaaa	120
acatgggaat	ggacgtctat	cgtttctcta	tctctgggtc	acgcataatt	cctaaggggt	180
cgccaagaca	cggaccagtc	aataaagtgg	gaatcgttta	ttacaataat	ttcatcaacg	240
agctgctcag	gaatgggtata	gagccttttg	tcacactgtt	tcactggggac	atgccacaag	300
ctctggaaga	tgagtacggg	ggattccgta	acaaaagagt	cgtggaggac	tttaacatat	360
ttgcagaagc	atgctttcga	gcctt				385

&lt;210&gt; 146

&lt;211&gt; 546

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 146

ctccccgtgc	cttttctctc	tcccttcatt	aattctctct	tccgagatct	gatttttccct	60
cacttttccc	agaaaataat	ccccccgatc	tccccccggg	aattcccccc	cggccgttcg	120
attccggcgc	gcgctccggc	gatcgctcgc	tcgctcgcta	gccggttctt	ctctcgctcg	180
ttccaccgga	gatggcgggc	gaatggatac	tgacgttgac	cgcgcagacg	ccgacgaaca	240
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gatttcagag	ttgtttgaga	gaaattcgag	cccgtgctac	tgacgttgag	aataaggaaa	480
aaggaaattaa	aatttcaaag	aaagattggg	agaaaactgca	cctccacatt	tctttcttta	540
catttc						546

&lt;210&gt; 147

&lt;211&gt; 786

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 147

tcactctcgg	gcattccgcc	agcacaccat	ctttcgtccg	tcattcaacc	tctatagatc	60
ggctctctcc	aggtacctgg	ttcgcttcc	ctgcatgttt	tttagaccaa	tagtttccc	120
acttacggaa	tttggttag	aattaggccc	tgcaaaagtt	ttatagcttc	ctctggggta	180
acggtagctt	acaggggtga	attcgttgga	gcacgcgtga	gaagctacag	acatgagtag	240
caatggcaac	gggcaaaaac	aaggaggggg	ctttttcgcc	gccttcgcct	cgggcctctc	300
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cacctccatg	gtgacacttc	cagtcattat	ctttgagcct	atgacgatgc	ttcagaagag	540
tgctgagtta	atggagtaca	cttatttgct	tgacatggca	gatgagtgtg	aagatcccta	600
tctcaagatg	gcttatgcag	catcatgggc	aatttctgtc	tatcctgcat	accagaggag	660
ttggaagccc	tttaacccta	ttcttggaga	aacttatgaa	atgggtcaatc	atggagggat	720
cacatttatc	gcagagcagg	tcagccacca	tcctccatgg	gctcagccta	tgccagaaat	780
gacatt						786

&lt;210&gt; 148

&lt;211&gt; 1748

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 148

ccgtcattca	acctctatag	atcggtcttc	tccaggggtg	aattcggttg	agcatcggtg	60
agaagctaca	gacatgagta	gcaatggcaa	cgggcaaaaa	caaggagggg	gctttttcgc	120
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ggaagctcat	cgaggtagat	ggcgaaaga	ggaccaggat	agttattgga	aaatgatgca	300

aaaatatatt	ggagcagatg	tcacctccat	ggtgacactt	ccagtcatta	tctttgagcc	360
tatgacgatg	cttcagaaga	gtgctgagtt	aatggagtag	acttatttgc	ttgacatggc	420
agatgagtgt	gaagatccct	atctcaagat	ggcttatgca	gcacatggg	caatttctgt	480
ctatcctgca	taccagagga	gttggaaagcc	ctttaaccct	attcttggag	aaacttatga	540
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agaggtatgg	agggctcgag	atcttccaaa	gaatgacaaa	tttcaatata	catattttgc	1080
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acttccgtgc	catgagttga	atgaaatttt	aaaagataat	ggaagctgcc	tccatgtgat	1680
cagaaacttg	gcatataatt	gatatcatgt	acctatgttt	tggtttgggg	aagaaaaaaa	1740
aaaaaaaa						1748

&lt;210&gt; 149

&lt;211&gt; 428

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 149

ccgtcattca	acctctatag	atcggctctc	tccagaatta	ggccctgcaa	aagttttata	60
gcttcctctg	gggtaacggt	agcttacagg	gttgaattcg	ttggagcatc	ggtgagaagc	120
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attggagcag	atgtcacctc	catggtgaca	cttcagtc	ttatctttga	gcctatgacg	420
atgcttca						428

&lt;210&gt; 150

&lt;211&gt; 419

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 150

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tctacatttc	tcttgccaac	cgcgtgagca	tcatcgggga	actaagacca	aagtcaaaag	419

&lt;210&gt; 151

&lt;211&gt; 401

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 151

cttcatctaa	agggcgcat	tgcaaaccct	tcaacccttt	actgggggaa	acttatgaag	60
ctgactatcc	ggagagagga	gttcacttct	tttccgagaa	ggttagtcac	caccctactc	120
tcattgcttg	ccattgagaa	ggaaggggtt	ggaaattctg	ggctgacagc	aatttaagga	180
caaaatcttg	gggacaatct	attcagcttg	atcctgtggg	agcacttacc	cttgagtttg	240
atgatggcga	gatttttcaa	tggaataagg	taacaactag	catcaacaat	cttatcattg	300
gaaaagttaa	ctgtgatcat	catggtgtca	tgaatataca	tggaaccac	caatattcat	360
gcaaattgaa	gttcaaggag	ccatctattc	ttgccgaact	c		401

&lt;210&gt; 152

&lt;211&gt; 349

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 152

cgcacatgcg	attggtctat	gcgggcatca	tgggctatat	cagtctatta	tgccatcaaa	60
aggacgtgga	agcctttcaa	tcctattctt	ggggagactt	atgaactggc	aaatcatggc	120
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&lt;210&gt; 153

&lt;211&gt; 533

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 153

ctctcgttta	cgtcctgata	gatatgcact	tgagccgggt	gaccttccta	aagctgggtg	60
tgaaaagagc	agcttggagg	aaaggcaaa	aggagagaaa	aagaaccgag	aaatgaaagg	120
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agcatgttat	tcgccccttc	gttctttctac	tcagcatttt	tttattcata	ggagatcgta	480
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&lt;210&gt; 154

&lt;211&gt; 354

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 154

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agtcgagcac	tacaaagaac	tgatggaaaa	atttcatcat	gtttcaacta	cctttttacg	180
gcttggaagg	ggatatcagg	aagcaattga	agaaataact	aagaagatgg	gtgctgggat	240
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tgctgcagga	ctagttggat	ttggtttgtc	acgactcttt	catgcagctc	agct	354

&lt;210&gt; 155

&lt;211&gt; 675

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 155

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agattcaggt	caggtcatta	attgcaggaa	tcggtacact	gccatggcaa	tctatacgcc	120
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gcgcagatcc	aacggattac	ggggaatata	gagcaaaagt	ttcgaaaagt	caacagatga	300

caatggcatt	gccatcgaag	ctgctggagg	aacggatggt	atcatcgtgg	gagcaggagt	360
cgcgggttcg	gctctggctt	acacacttgg	caaggatgga	agacgtatac	atgtaattga	420
gagagacttg	agtgagcctg	accggattgt	aggggaactt	ttacagccag	gtggatattt	480
gaaattgatt	gagctggggac	ttcaagattg	tggtgaagga	attgatgccc	agagtatatt	540
tggggatgct	ttattcaagg	aaggaaaaga	tactaaagtg	gcataatccgt	tagaaaacca	600
ccatgcagat	agagctggaa	ggagtttcca	caatggacgc	ttcatccagc	gcatgcggga	660
aaaggctgct	tcact					675

&lt;210&gt; 156

&lt;211&gt; 373

&lt;212&gt; DNA

&lt;213&gt; Pinus radiata

&lt;400&gt; 156

tgccttacga	cagattcagg	tcaggtcatt	aattgcagga	atcgggtacac	tgccatggca	60
atctatacgc	ctcaaccagc	acatcgactg	atatcgtggt	ctacaatgga	gaatcatact	120
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ggagcaggag	tcgcgggttc	ggctctggct	tacacacttg	gcaaggatgg	aagacgtata	360
catgtaattg	aga					373

&lt;210&gt; 157

&lt;211&gt; 522

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 157

cgcaaacata	gtctgagctt	gcacatttcg	gaaatcccgt	aaagcagcac	agcttgcacc	60
gcaagagccg	agctccatcg	gcggaagctt	ctcgtatctc	ctgctgtcgc	gtggcggagg	120
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tcttggtctc	gttcctgggg	atcttcctgc	tgtacaaggg	gctcgggaag	cagaagagga	240
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tgcccgccgg	cgctcgatggg	agcaccgaca	tcgtcattgt	cggagcaggg	gtcgcgggtg	360
cggctctcgc	ttacgccctc	gggaaggatg	gacgtcgcgt	gcgtgtaatt	gagagggacc	420
tgacggagca	agatagaatt	gtcggcgagc	ttcttcaacc	aggagggtac	ctgaaattga	480
tgaatttga	ccttgcagat	tgcgtgcaaa	caattgatgc	cc		522

&lt;210&gt; 158

&lt;211&gt; 898

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 158

ctcgggtcga	agtataaacc	tcaggaagaa	tttgttgaat	ggattcaaaa	gggaacaaaa	60
cctatatata	tcgggttttg	gagcatgcct	cttgaagatc	ccaagaaaac	tacagatata	120
ataattaagg	cattaacgga	taccggacaa	agagggatag	ttggtcgagg	ttgggggtgat	180
cttgggaccc	ttctggatgt	tccagacagt	gttttccttt	tggaggattg	tccgcgatgat	240
tggctgtttc	cccaatgttc	agctgtggtt	catcacgggtg	gtgctggaac	aacagctaca	300
ggactaaaa	cagggtgtcc	tacaaccata	gttcctttct	ttggggatca	gttcttcttg	360
ggcgatagg	tccaccaaag	aggccttggc	cctgcaccaa	taccaatctc	ccagctcagc	420
gtcgagaacc	tttcagatgc	cataagattt	atgcttcaac	ctgaggtaaa	gtctcaggca	480
atggaaatgg	cgaagctgat	agaaaatgag	gatgggggtg	cggctgctgt	cgacgcgttt	540
catcggcatc	tgcgggaaga	gtttccctcg	tcgagtgtgt	cctcggacgg	tgaggagcac	600
cccaaccctt	tcctgtggct	cttcctccaa	gttgagaagt	ggtgctgcct	gccatgtagt	660
aaataggggc	cttccttttg	ataaattgag	agtgggtgta	tagaagtgat	agatgtcttc	720
cttattgttt	tctgtccctc	agttaccatt	tttttttctt	ttcaaatttg	tttcaaatca	780
ttcatttcat	tcttatcagg	gtttggctga	ccattgtatt	cagcatagca	taagatttaa	840
ttttgccact	gcttcttgtg	taaaatcact	aggcttcatt	tggaactgtt	atatttttg	898

&lt;210&gt; 159

&lt;211&gt; 342

&lt;212&gt; DNA

<213> *Eucalyptus grandis*

&lt;400&gt; 159

ctcgataatt	gccctcatga	ctggcttttc	ctgcgctgca	gtgctgtggt	acatcatgga	60
ggagctggta	caaccgctgc	cggctcttaa	gctgcgtgtc	caacaacagt	tgtacctttc	120
tttggggatc	agcccttttg	gggagaacgg	gtgcatgcaa	gggggggtgg	cccagtgcca	180
attccagttg	atgaattttc	tcttgaaaag	ttggttgatg	caatacgttt	catgcttgat	240
ccaaagggtg	aacagtgtgc	agaagaacta	gccaaagaca	tggaacatga	agatggagtg	300
gagggagcag	tgaaggcttt	ctacaaacac	tttcacgcg	aa		342

&lt;210&gt; 160

&lt;211&gt; 582

&lt;212&gt; DNA

<213> *Pinus radiata*

&lt;400&gt; 160

atgcttgctg	tgccaaaaac	cgatgtaatt	tattattgtg	cgcagggatt	ccttctgctc	60
cttatgatcc	cctaaccctt	aaatcgtagc	agtgaagcca	ttaacgattt	ttgcgggttc	120
agaaagattc	actgaatcgc	ttactaaaac	tctgtttcag	gaatggcaac	aggaggagga	180
gcgttggtac	tggcctcagg	aatgggaggc	aacattgaga	aagaacaaat	gctgaccgct	240
gttgaagagt	acgaaaaata	tcacatgtac	tatggtgggtg	atgaaggctc	gagaaaatct	300
aactatacag	acatggtaaa	ttaatactat	gatctggcga	ctagtttcta	tgagtatgga	360
tggggggagt	cttttcattt	tgctcacaga	tggaaagggg	agaccctccg	agaaagtata	420
aagcgccatg	aacattttct	tgctcttcac	ctttgtttta	agcctgcaat	gaagggtattg	480
gatgttggtg	gtgggattgg	aggtccactg	agagaaattg	ctagggttcag	tcggacttcg	540
atcacaggat	tgaataataa	tgcatatcag	atatcaagag	ga		582

&lt;210&gt; 161

&lt;211&gt; 552

&lt;212&gt; DNA

<213> *Eucalyptus grandis*

&lt;400&gt; 161

cttcttgctt	gtctctgcct	ctctctctct	cgttcctagg	gttctgaagc	tgatcctcct	60
cctgcattgt	cctcattctg	ggcgggggtg	ccacaatgtc	gaaagcagga	gcgatggatc	120
tggcgacggg	ccttggcggg	aagatggaca	agagcgacgt	cctgtccgcc	gttgacaagt	180
atgagaagta	tcattgtctg	tatggaggtg	atgaggaaga	aaggagagct	aactatagtg	240
acatggtgaa	taaatattat	gatcttgcta	ccagctttta	tgagttcggc	tggggagaaat	300
ctttccattt	tgcccacaga	tggaaagggg	agtctctacg	agagagcatt	aagagacatg	360
aacactttct	tgcattacag	ctaggcttaa	aacctgggca	caagggtgctg	gatgtcgggt	420
gcggaattgg	tggaccgctt	agggaaatag	ctcgattcag	ctccgcattct	gttacaggat	480
taaacaacaa	tgagtaccag	ataacaaggg	gaaaggaact	aaaccgcatt	gcaggcggtg	540
acaagacatg	cg					552

&lt;210&gt; 162

&lt;211&gt; 401

&lt;212&gt; DNA

<213> *Eucalyptus grandis*

&lt;400&gt; 162

cttcttcttg	cctgtctctg	cctctctctc	tctctcgttc	ctagggttct	gaagctgac	60
ctcctcctgc	attgtcctca	ttctgggcgg	ggtggccaca	atgtcgaaag	caggagcgat	120
ggatctggcg	acgggccttg	gcggaagat	ggacaagagc	gacgtcctgt	ccgccgttga	180
caagtatgag	aagtatcatg	tctgctatgg	aggtgatgag	gaagaaagga	gagctaacta	240
tagtgacatg	gtgaataaat	attatgatct	tgctaccagc	ttttatgagt	tcggctgggg	300
agaatctttc	cattttgccc	acagatggaa	aggggagctc	ctacgagaga	gcattaagag	360
acatgaacac	tttcttgcat	tacagctagg	cttaaacct	g		401

&lt;210&gt; 163

&lt;211&gt; 446

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 163

ggcaatgcc a tggactgtg ctctccaac actttccgaa tacatgggtg aaaatggatg 60  
 gacgaaatgc ttttcaagaa taagcgatgt tggttggctt gcttacctag tgtactgtc 120  
 aatatatctt gtaatggcgg agtttgggat atattggatg cacagagagc tgcagacat 180  
 taaacccctt taatggcgg taatggcgg taatggcgg taatggcgg taatggcgg 240  
 ttctctttt gccggcttg cctttgacg tctagacgg atactgcagg cggtgccaca 300  
 tggtaggca ttattctctg tggcagc ttttccatg ttttccatg ttttctttct 360  
 cgaggccata tggacagcaa atatccatga ctgcatccat ggtaagcttt ggctgtgat 420  
 gggcgctggt tatcacacca tccacc 446

Y2K

&lt;210&gt; 164

&lt;211&gt; 823

&lt;212&gt; DNA

&lt;213&gt; Eucalyptus grandis

## BUSINESS CONTINUITY PLAN

&lt;400&gt; 164

gttcatgtca tgcctgccca acatgattgt catggcgcca tctgatgaag atgagctagt 60  
 ggacatggta gaaactgctg ccatagttga tgatcgctca atttgcttcc gctatccacg 120  
 ggggtgcaatt gttcggacag acaagtctct cagtcaaggg atccccattg agattggaaa 180  
 ggggagaatt cttgcagaag gtaaagatgt tgggttctt ggttatgggt caatggttca 240  
 gaactgtgtc aaagctcggg ccttcttttc aaagctcggg attgaggtga cagtcgctga 300  
 tgcaagattc tgcaaacccg ttgacatagg attacttagg gaggttgtg aaaatcatgc 360  
 cttctgtgct actgttgagg aaggatccat cggagggttc gggteccatg ttgcacagtt 420  
 cattgcactt gatggacggc ttgatgggag aataaagtgg cggccgattg ttttacctga 480  
 tgcctatgta tgcctatgta tgcctatgta tgcctatgta tgcctatgta tgcctatgta 540  
 tcacattgcc gcaactgtgt tgagctctct tggcgggaca cgcgaagctc ttctgttgat 600  
 gtgctagggt cctcgaatt cttccgccc ttttccatg gaaatgggct cgtgctgatg 660  
 ccgcagtact gataagccag acatgttaat gaagcttgag caaagatggc ttactcgccg 720  
 actaccatgt gtccagaatg ctgtgttgat tgttggcatg cagtacgttc tatcgccaga 780  
 atgcagaact catttctgag aagcttattc ggagatgttt ctg 823

DOCUMENT AND INFORMATION SERVICE  
CENTRE

&lt;211&gt; 90

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

17 TOOP ST

&lt;400&gt; 165

Phe Asp Asp Val Ser Ala Leu Phe His Asp Leu Asp Asn Gly Met Leu  
 1 5 SEAVIEW 15  
 Pro Ile Ser Val Ile Phe Pro Tyr Leu Pro Ile Pro Ala His His Arg  
 20 25 LOWER HUTT 30  
 Arg Asp Lys Ala Arg Lys Lys Leu Ser Glu Ile Phe Ala Asn Ile Ile  
 35 40 45  
 Ser Ser Arg Lys Cys Ala Gly Lys Ser Glu Glu Asp Met Leu Gln Cys  
 50 55 60  
 Phe Ile Asp Ser Lys Tyr Lys Asn Gly Arg Pro Thr Thr Glu Ala Glu  
 65 70 75 80  
 Val Thr Gly Leu Leu Ile Ala Ala Leu Phe  
 85 90

&lt;210&gt; 166

&lt;211&gt; 40

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

PREPARED DECEMBER 1999 REVISION 7

&lt;400&gt; 166

Tyr Leu Leu Thr Asn Lys Lys Tyr Leu Ser Ala Val Ser Asn Glu Gln

1	5	10	15
Lys His Leu Met Glu Lys His Gly Asn Val Asp His Asp Val Leu			
20	25	30	
Ser Glu Met Asp Val Leu Tyr Arg			
35			

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<400>	167
Arg Leu Val Phe Ala Glu Gly Glu Asp Gly Pro Tyr Leu Tyr Ser Thr	
1	5
Asn Arg Val Phe Ile Trp Glu Phe Asp Pro Glu Ala Gly	
20	25
Thr Ala Glu Arg Ala Glu Val Glu Ala Ala Arg Gln His Phe Tyr	
35	40
Asp His Arg His Gln Val Lys Pro Cys Gly Asp Leu Leu Trp Arg Met	
50	55
Gln Phe Leu Arg Glu Lys Glu Phe Lys Gln Thr Ile Pro Pro Val Arg	
65	70
Val Glu Asp Gly Glu Ile Thr Tyr Asp Lys Ala Ser Thr Ala Leu	
75	80
Lys Arg Ala Val His Phe Phe Ser Ala Leu Gln Ala Ser Asp Gly His	
105	110
Trp Pro Ala Glu Asn Ala Gly Pro Leu Phe Phe Leu Pro Pro Leu Val	
120	125
Met Cys Val Tyr Ile Thr Gly His Leu Asp Ala Val Phe Pro Ala Glu	
135	140
His Arg Lys Glu Ile Leu Arg Tyr Ile Tyr Asn His Gln Asn Glu Asp	
145	150
Gly Gly Trp Gly Leu His Ile	
155	160
4.2.	Pre 24 December 1999
<210>	168
<211>	Post 1 January 2000
<212>	PRT
5.	STAFFING / SUPPORT
<213>	Eucalyptus grandis
6.	INVOKING THE PLAN
Met Asp Asp Ile Val Ser His Glu Phe Glu Gln Lys Arg Gly His Val	
1	10
Val Ser Ala Val Glu Leu Leu Ile Lys Tyr Arg Gly Val Ser Glu Gln	
25	30
Glu Ala Val Glu Leu Gln Lys Arg Val Ile Asp Ala Trp Lys Asp	
40	45
Thr Asn Glu Glu Phe Leu Arg Pro Ile Ala Val Pro Met Pro Ile Leu	
50	55
Thr Arg Val Leu Asn Leu Ser Arg Val Ile Asp Val Leu Tyr Ser Asp	
65	70
Gly Asp Asn Tyr Thr His Ser Glu Thr	
75	80
9.	COMMUNICATION PLAN
85	
10.	APPENDIX
<210>	169
<211>	Contact Details
<212>	PRT
<213>	Telephone Tree Details
11.	REPORT OF TESTING
Met Glu Asp Asp Arg Asp Arg Gly Leu Leu Tyr Asp Ser Asp Pro Pro	

1	5	10	15
Ser	Pro	Ser	Leu
20	25	30	
Thr	Phe	Phe	Asp
35	40	45	
His	Met	Leu	Pro
50	55	60	
Leu	Ala	Val	Thr
65	70	75	
Arg	Glu	Arg	His
80	85	90	
His	Met	Leu	Pro
95	100	105	
His	Met	Leu	Pro
110	115	120	
His	Met	Leu	Pro
125	130	135	
His	Met	Leu	Pro
140	145	150	
His	Met	Leu	Pro
155	160	165	
His	Met	Leu	Pro
170	175	180	
His	Met	Leu	Pro
185	190	195	
His	Met	Leu	Pro
200	205	210	
His	Met	Leu	Pro
215	220	225	
His	Met	Leu	Pro
230	235	240	
His	Met	Leu	Pro
245	250	255	
His	Met	Leu	Pro
260	265	270	
His	Met	Leu	Pro
275	280	285	
His	Met	Leu	Pro
290	295	300	
His	Met	Leu	Pro
305	310	315	
His	Met	Leu	Pro
320	325	330	
His	Met	Leu	Pro
335	340	345	
His	Met	Leu	Pro
350	355	360	
His	Met	Leu	Pro
365	370	375	
His	Met	Leu	Pro
380	385	390	
His	Met	Leu	Pro
395	400	405	
His	Met	Leu	Pro
410	415	420	
His	Met	Leu	Pro
425	430	435	
His	Met	Leu	Pro
440	445	450	
His	Met	Leu	Pro
455	460	465	
His	Met	Leu	Pro
470	475	480	
His	Met	Leu	Pro
485	490	495	
His	Met	Leu	Pro
500	505	510	
His	Met	Leu	Pro
515	520	525	
His	Met	Leu	Pro
530	535	540	
His	Met	Leu	Pro
545	550	555	
His	Met	Leu	Pro
560	565	570	
His	Met	Leu	Pro
575	580	585	
His	Met	Leu	Pro
590	595	600	
His	Met	Leu	Pro
605	610	615	
His	Met	Leu	Pro
620	625	630	
His	Met	Leu	Pro
635	640	645	
His	Met	Leu	Pro
650	655	660	
His	Met	Leu	Pro
665	670	675	
His	Met	Leu	Pro
680	685	690	
His	Met	Leu	Pro
695	700	705	
His	Met	Leu	Pro
710	715	720	
His	Met	Leu	Pro
725	730	735	
His	Met	Leu	Pro
740	745	750	
His	Met	Leu	Pro
755	760	765	
His	Met	Leu	Pro
770	775	780	
His	Met	Leu	Pro
785	790	795	
His	Met	Leu	Pro
800	805	810	
His	Met	Leu	Pro
815	820	825	
His	Met	Leu	Pro
830	835	840	
His	Met	Leu	Pro
845	850	855	
His	Met	Leu	Pro
860	865	870	
His	Met	Leu	Pro
875	880	885	
His	Met	Leu	Pro
890	895	900	
His	Met	Leu	Pro
905	910	915	
His	Met	Leu	Pro
920	925	930	
His	Met	Leu	Pro
935	940	945	
His	Met	Leu	Pro
950	955	960	
His	Met	Leu	Pro
965	970	975	
His	Met	Leu	Pro
980	985	990	
His	Met	Leu	Pro
995	1000	1005	
His	Met	Leu	Pro



130

## Media Contacts

~~211-83~~

<210> 174

<212> PRT

## Accidents and Mo

His	Ser	Leu	Gln	Gln	Lys	Glu	Leu	Lys	Gln	Leu	Ser	Arg	Trp	Trp	Lys
50					55					60					
Asp	Ser	Gly	Phe	Ser	Gln	Leu	Thr	Phe	Thr	Arg	His	Arg	His	Val	Glu
65				70						75				80	
Phe	Tyr	Thr	Leu	Ala	Ser	Cys	Ile	Ala	Thr	Glu	Pro	Lys	His	Ser	Ala
			85						90					95	
Phe	Arg	Leu	Gly	Phe	Ala	Lys	Thr	Cys	Tyr	Leu	Gly	Ile	Val	Leu	Asp
			100						105					110	
Asp	Ile	Tyr	Asp	Thr	Phe	Gly	Thr	Met	Glu	Glu	Leu	Glu	Leu	Phe	Thr
			115						120					125	
Ala	Ser	Arg	Trp	Asp	Ser	Ala	Arg	Glu							
			130						135					140	
Scenario Presented															
Invoke Plan Decision															
No access to Seaview area															
No building access															
Building access, no utilities															
Building access, no IT															
Building access, no IPOL															
No Mail Delivery															

<400> 175  
 Leu 3.1 No Access to Seaview  
 Thr Asn Phe Pro Arg Leu Glu Ala Arg Asn Tyr Ile Asp Val Phe  
 1 5 10 15  
 Gly In the event that Seaview suffers major infrastructure problems the Impact Assessment Team  
 will:  
 20 25 30  
 Thr Glu Contact staff by phone tree  
 35  
 • Leave recorded phone message on (04) 568 0744 for staff information and (04) 568 0720 for  
 Ser Leu Gln Gln Lys Glu Leu Lys Gln Leu Ser Arg Trp Trp Lys Asp  
 50 Clients information. 55 60  
 Ser Gly Phe Ser Arg Leu Thr Phe Thr Arg His Arg His Val Glu Phe  
 65 70 75 80  
 Tyr Thr Leu Ala Cys Cys Ile Ala Thr Glu Pro Lys His Ser Ala Phe  
 85 90 95  
 Arg In the event the Documents & Information Service Centre remains closed the Impact Assessment  
 Team will: 100 105 110  
 Ile Tyr Asn Thr Phe Gly Thr Met Glu Glu Leu Glu Leu Phe Thr Ala  
 115  
 • Leave recorded phone message on (04) 568 0744 for staff information and (04) 568 0720 for  
 Ala Ile Lys Arg Trp Asp Pro Ser Ala Arg Glu Cys Leu Pro Glu Tyr  
 130 Clients information. 135 140  
 Met Lys Gly Ile Tyr Met Val Phe Tyr Asp Ala Leu Ile Lys Trp Leu  
 145 150 155 160  
 Glu 3.2 Building Access no Utilities

If the building is open but there is no power, water or sewerage the Impact Assessment Team  
 will: <210> 176

- If staff not present, contact by phone tree.
- If staff already present, consider sending staff home.
- If no power, the leader will contact Manager IT Operations.
- Leave recorded phone message on (04) 568 0744 for staff information and (04) 568 0720 for

Trp Ser Clients' Information Phe Val Pro Ser Phe His Glu Tyr Ile Ala Thr  
 1 5 10 15  
 Ala Ser Ile Ser Val Ser Gly Pro Thr Leu Ile Leu Ile Cys Val Leu  
 20 25 30  
 Phe Thr Gly Glu Leu Leu Thr Asp His Ile Leu Cys Gln Ile Asp Tyr  
 35  
 Arg If all services are available except IT systems the Impact Assessment Team will:  
 40  
 • Re-assign staff to other work. 50  
 Asp Thr Lys Thr Tyr Gln Ala Glu Arg Gly  
 65 70

### 3.5 Building Access but no IPOL

<210> 177

If the IPOL system is not available the Impact Assessment Team will:

- Re-assign staff to other work.

<213> Eucalyptus grandis

Leu 1 3 6 10 15  
Ala Asp Leu Lys Gly Glu Phe Leu Asn Arg

Ser Ser Gln Leu Phe Cys Met Glu Asn Asp Gly Phe Thr His Ser His  
35 40 45

~~Glu Thr Glu Ile Lys Glu His Val Lys Lys Ile Leu Phe Glu Pro Val~~  
50 55 60 PLAN SUMM 60

#### 4. PLAN SUMMARY<sup>60</sup>

<210> 178

<213> Eucalyptus grandis

<400> 178

1	Task	5	Responsibility	Target Date	Completed
Val	Seek assurance from suppliers for Y2K compliance	13	Sue Whiteman	31 July 1999	Yes
Met	Test all software and hardware for Y2K compliance	40	Shirley Herewin	13 July 1999	Yes
Met	Discuss with Support Services arrangements for alternative accommodation	55	Shirley Herewin	30 September 1999	Yes
65	Contact key suppliers to obtain and confirm contact names	70	Shirley Herewin	31 October 1999	Yes
Cys	Distribute copies of BCP plan to all Staff for home and work	105	Sue Whiteman	15 November 1999	Yes
Gly	Inform Clients and staff of Holiday Period telephone number on which messages will be left	125	Glanet Dobber	10 December 1999	Yes
145	Arrange necessary stationery for date stamping	135	Shirley Herewin	15 December 1999	Yes
Arg	Arrange storage for mail and cheques	150	Shirley Herewin	15 December 1999	Yes
	Arrange van to be full of petrol	160	Shirley Herewin	24 December 1999	

Phe 431 Arg After 1 January 2000 Ala Arg Leu Gln Ser Ile Lys Cys Ala  
 195 200 205

Task	Gly	Lys	Asn	Leu	Tyr	Met	Responsibility	Cys	Ser	Target Date	Asp
Check that location is accessible	215						Armourguard	220		12:30 am	Armourguard
Met: Gly Met							Lys Gly Val	Gln	Asn	Security to contact Neville	
building secure, power on	230							235		240	Harris (Diane) minus as backup
							Met Asp Val	Leu	Gly	to Neville	
								250		255	
If no power or access to office	245						Neville Harris	Asn	Trp	By 1 am	
Met: Gly Met							260			270	Jan 6 am
contact Mike	260						Alonso	Lys	Gly	Asp	Val Val
							Alonso	Lys	Gly	Asp	Val Val
BRB testing of IT LAN/WAN	280						Michael Brosnahan	285		Jan 1 6am (initial IT ck)	Joe
Met: Gly Met							Joe Flynn (can Michael	Gln	Val	Flynn report to Kathryn	
systems	290						check)	300		McInteer 11am 1 Jan	
							Met Ala Gly	Ala	Leu	Full BRB test start by 1pm 1	
								315		320	Jan. MB to report to Kathryn
										McInteer by 3pm	

Asn Ala His Ala Ser Asn Ile Val Ala Ala Ile Phe Ile Ala Thr Gly	325	330	335
Glrs of Applications	340	345	350
Ala Ile Asn Asp Gly Lys Asp	355	360	365
Val Glu Val Gly Thr Val Gly	370	375	380
Ala Cys Leu Asn Leu Leu Gly	385	390	395
Gly Ala Asn Ser Arg Leu Leu	405	410	415
Ala Ala Glu Leu Ser Leu Met	420	425	430
Val Office site assessment	435	440	445
Complete checklist from Ministry	450	455	460
IT Reboots office servers	465	470	475
Testing of Applications and processing and Application sponsor report back to Michael Brosnahan with sign off PRT	480	485	490
<213> Eucalyptus grandis	495	500	505
<400> 179	510	515	520
Ser Arg Asn Arg Glu Ala Asp Ala	525	530	535
Glrs Ser Ala Cys Leu Asn Leu Leu	540	545	550
Ala Gly Ala Asn Ser Arg Leu Leu	555	560	565
Leu Ala Ala Glu Leu Ser Leu Met	570	575	580
Val Office site assessment Complete checklist from Ministry 1pm 1 Jan.	585	590	595
Val Ser Ser	600	605	610
Conduct post disaster audit appraisal and document results. Amend plan where necessary	615	620	625
<b>Implement Below Only If Problem Identified</b>			
<213> Eucalyptus grandis	630	635	640
Check outcome of testing on 1 January 2000 and prepare Recovery Team to mobilise if necessary	645	650	655
Leu Tyr Met Gly Phe Ser Lys Ser	660	665	670
1 Invoke Telephone tree	675	680	685
Met Inform any personnel of situation and invoke appropriate action plans	690	695	700
Phe Carry out PPA	705	710	715
Monitor situation	720	725	730
Lys Pys PLO Ala Ala Val Asn Tip	735	740	745
Conduct post disaster audit appraisal and document results. Amend plan where necessary	750	755	760
Val Cys Glu Ala Val Ile Lys Gly	765	770	775
65	780	785	790

&lt;210&gt; 181

&lt;211&gt; 81

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 181

Ser Ile Lys Cys Ala Ile Ala Gly Lys Asn Leu Tyr Leu Arg Phe Ser  
 1 5 10 15  
 Cys Ser Thr Gly Asp Ala Met Gly Met Asn Met Ile Ser Lys Gly Val  
 20 25 30  
 Gln Asn Val Met Asp Phe Leu Gln Lys Asp Phe Pro Asp Met Asp Val  
 35 40  
 Met Gly Ile Ser Gly Asn Phe Cys Ser Asp Lys Lys Pro Ala Ala Val  
 50 55 60  
 Asn Trp Ile Glu Gly Arg Gly Lys Ser Val Val Cys Glu Ala Val Ile  
 65  
 Lys The Impact Assessment Team is required to be on call to assess the situation after Computer  
 Services BRB IT have completed testing on 2 January 2000. Leader, Janet Dobbie will be  
 responsible for informing the rest of the Team.

Those on call on 1 January 2000 need only be contacted for a progress report on the outcome of  
 IT services availability. They will be required to make preparations to return to Wellington if the  
 need arises. PRT

<213> Pinus radiata

Person	Availability	Date & Time	Back-up Resource
Janet Dobbie	On call	Glu Glu Val Val Lys	Diane Imus
Sue Whiteman	On call	10 15	
Gary Jones	On call	Glu Leu Asn Met Leu	
Shirley Herepuni	On call	25 30	
Rest of DISC staff	On site unless otherwise advised	15 January 8am	Phe Asn
35	40	45	

Ala His Ala Ser Asn Ile Val Ser Ala Ile Tyr Ile Ala Thr Gly Gln  
 50 55 60

Asp Pro Ala Gln Asn Val Glu Ser Ile Thr Met Met Glu  
 65 70 75 80

Ala Val Asn Glu Gly Arg Asp Leu His Ile Ser Val Thr Met Pro Ser  
 85 90 95

Ile Glu Val Thr Ile Val Gly Gly Thr Gln Leu Ala Ser Gln Ser  
 100 105 110

Ala The Impact Assessment Team are responsible for determining the operational status of DISC  
 and the impact on IPONZ operations 125

Gly Ala Asn Ala Arg Leu Leu Ala Thr Ile Val

On or by 2 January 2000 Janet Dobbie will check:

- Power is OK (Armourguard to report back to Neville).
- Access to IPOL is available - Mike Brosnahan by 4pm 1 Jan
- If IPOL or IT is not available BRB IT will be called to assess the problem and make an assessment call on how to proceed.
- If all IT and IPOL OK - Diane Imus to contact Janet and keep informed.
- If the system is still unavailable at 5.30pm on 4 January 2000, Janet Dobbie will inform

Met Asp Val Arg Arg Arg Gln Lys Ser Leu Arg Leu Pro Ala Pro Gly Val  
 1 5 10 15  
 Impact Assessment Team and staff using the telephone call free.

Asp Pro Arg Arg Arg Gln Lys Ser Leu Arg Leu Pro Ala Pro Gly Val  
 20 25 30  
 As a result the Impact Assessment Team will determine the effects of any problems on

Asp Arg Arg Arg Arg Gln Lys Ser Leu Arg Leu Pro Ala Pro Gly Val  
 35 40 45  
 • 3 Resources

Pro Leu Pro Leu Tyr Leu Thr Asn Ala Val Phe Phe Thr Leu Phe Phe  
 50 55 60  
 • Decide on the appropriate courses of action for recovery purposes and mobilise the

Ser Val Arg Arg Arg Arg Gln Lys Ser Leu Arg Leu Pro Ala Pro Gly Val  
 65 70 75 80  
 Recovery Team

Ser Val Pro Leader - Notify staff and clients the recorded message on phones. Ile Val  
 85 90 95  
 • Leader - Notify BRB Impact team.

Ser Leu Ile Ala Ser Phe Ile Tyr Leu Leu Gly Phe Phe Gly Ile Gly  
 100 105 110

Phe Val Gln Phe Phe Ile Ala Ser His Asp Ala Trp Asp Val  
 115 120 125  
 6.2 Implement a manual system of date stamping

Leu Asp Asp Glu Val Ala Val Gly Gly Asp Gly Phe Leu Pro Glu Asp  
 If a manual system is required the Team will:

PAGE 9

&lt;210&gt; 185

&lt;211&gt; 194

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis COMMUNICATION PLAN

Sue Whiteman will be responsible for ensuring copies of the plan are distributed via e-mail to all staff.

Met Val Arg Arg Arg Pro Pro Lys Pro Pro Leu Pro Ser Ala Ala  
1 5 10 15

Arg Gly Gly Gly Arg Gly Pro Ala Ser Ser Ser Pro Pro Leu Glu Pro  
20 25 30

Pro Lys Ala Ser Asp Ala Leu Pro Leu Pro Leu Tyr Leu Thr Asn Ala  
A further copy will be distributed to all staff on the 22nd December 1999.

Val Phe Phe Thr Leu Phe Phe Ser Val Ala Tyr Tyr Leu Leu His Arg  
Communicating to staff of developments - phone message on a particular number (04 568 0744)

Trp that they may use. Phone message will be updated by Janet Dobbie. Thr Leu  
65 70 75 80

Pro Communicate to clients and BRB, MOU phone message (04 568 0720) to be changed by  
Janet Dobbie. 85 90 95

Leu Gly Phe Gly Ile Asp Phe Val Gln Thr Phe Ile Ala Arg Ala  
100 105 110

Poster on door if the building is not in use.

Ser His Asp Ala Trp Glu Asp Leu Asp Asp Asp Val Asn Arg Gly Phe  
115 120 125

Gly Media release (if necessary) to be done by the Ministry Communications Unit ONLY  
130 135 140

Pro Val Ile Ser Ala Leu Ser Ser Ala Glu Asp Glu Glu Ile Val Lys  
145 150 155 160

Ser Val Val Asp Gly Thr Ile Pro Ser Tyr Ser Leu Glu Ser Lys Leu  
165 170 175

10. APPENDIX

Name	Position	Contact Number
Janet Dobbie 138 Cockayne Road Ngairi Wellington	Manager, Document & Information Service Centre	(04) 568 0720 work (04) 479 7539 home 021 362 898 mobile
Gary Jones 28 Exploration Way Whitby	Team Leader Post Acceptance	(04) 568 0726 work (04) 234 1400 home
Shirley Pieroni P.O. Box 10 Shelly Bay	Team Leader Records Ser Leu Pro Ser Asp Phe	(04) 568 0731 work (04) 380 9028 home 025 405 957 mobile
Sue Whiteman State Highway 1 McKays Crossing, Paekakariki	Support Services Duty Manager for Building	(04) 568 0744 work (04) 292 8018 home 025 411 812 mobile
Diane Lewis 386c Karori Road Wellington	National Manager Corporate Services Business & Logistics Branch	(04) 470 2514 work (04) 476 3459 home 021 342 604 mobile
Janet Campbell SUNEX Ltd Debbie Monahan	Support Services Manager IPONZ Manager IPONZ	(04) 560 1601 work (04) 987 0708 home (04) 560 1615 work 021 306 098 mobile
Michael Grossman Trp Met Val Asp Ser Val	Operations, BRB Glu	(09) 913 4221 025 443 702 mobile
Glas McKenzie	BRB IT System Administrator	021 532 689 mobile
Nedax Security	Mark Eden	(04) 471 2836
Armourguard Security	Monitoring Centre	(04) 478 1226
DX<211> 140		Phone: (04) 473 9510
Post & Waste Couriers		Phone: (04) 499 2121
ASB Cleaners	Eric Reille	(04) 564 3249

&lt;213&gt; Pinus radiata

<400> 187		025 454 162
Met	BRB Corporate Phone Free (in contact order)	Ser Lys Leu Cys Leu Cys
1	Name 5	Address 10
Arg	511 481 Phe Gly Phe Ser	845 Gly Leu Lys Ala Ile
	Diane Innes 20	385C Karori Rd
Val	Pro Asn Leu Gly Met Cys	Karori, Wellington
	35	40
Met	Neville Harris	48 Ponsonby
	50	55
Arg	Arg Ile Ala Gly His His	Ser Asn Leu Trp Asp Asp
65	Andrew Bridgman 70	3 Fettes Cres 75
Ala	Ser Leu Ser Thr Ser Tyr	Seatons Heights
	85	90
Asp	Adam Feeley	36 Fortification Rd
	50	55
	Lys Leu Ile Gly Glu Val	58A Asn Ile Phe Asp Leu
	100	105
Glu	Rodney Grindley	26 Taupo Cres
	115	120
Trp	Karina Bach	102 Heke Street
	130	135
		140
		145
		150
		155
		160
		165
		170
		175
		180
		185
		190
		195
		200
		205
		210
		215
		220
		225
		230
		235
		240
		245
		250
		255
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		265
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		275
		280
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		385
		390
		395
		400

&lt;210&gt; 188

&lt;211&gt; 68

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 188

Leu Gly Met Pro Arg Arg Trp Lys Phe Ala Arg Pro Ser Met Ser Leu  
 1 5 10 15  
 Ser Thr Val Ala Ser Asp Asp Asp Ile Gln Arg Arg Thr Gly Gly Tyr  
 20 25 30  
 His Ser Asn Leu Trp Asn Asp Asp Val Ile Gln Phe Leu Ser Thr Pro  
 35 40 45  
 Tyr Gly Glu Leu Ala Tyr Arg Glu Arg Ala Glu Arg Leu Ile Asp Glu  
 50 55 60  
 Val Arg Asn Ile  
 65

&lt;210&gt; 189

&lt;211&gt; 99

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 189

Asp Asp Ala Val Ile Arg Arg Arg Gly Asp Tyr His Ser Asn Ile Trp  
 1 5 10 15  
 Asp Tyr Asp Phe Ile Gln Ser Leu Ser Ala Pro Tyr Gly Glu Pro Ser  
 20 25 30  
 Tyr Leu Glu Arg Ala Glu Arg Leu Ile Glu Glu Val Lys Lys Val Phe  
 35 40 45  
 Asn Ser Met Ser Glu Glu Asn Gly Glu Leu Ile Thr Pro Leu Asn Asp  
 50 55 60  
 Leu Ile Gln Arg Leu Trp Met Val Asp Ser Val Glu Arg Leu Gly Ile  
 65 70 75 80  
 Asp Arg His Phe Glu Asn Glu Ile Glu Ser Ala Leu Asp Tyr Val Tyr  
 85 90 95  
 Ser Tyr Trp

&lt;210&gt; 190





Ser Asn Ala Asn Gln Leu Ser Ser Met Gly Phe Ala Phe Ser Ser Gly  
 20 25 30  
 Ser Leu Tyr His Gln Val Met Arg Thr Lys Leu Gln Ser Met 175  
 To change your Voice Mail greeting from an external number phone (07) 560 1679  
 • Enter your mailbox number 35 40 45  
 Gly Arg Val Gly Lys Ala Tyr Ala Ser Ala Leu Ser Asp Gln Gly  
 • Enter your password number 50 55 60  
 Gln Tyr Ser Arg Lys Pro Thr Pro Leu Leu Asp Thr Ile Asn  
 65 • For an external greeting press 1 and then 5 to start recording your greeting. 80  
 Tyr Pro Ile His Met Lys Asn Leu Ser Ile Arg Gln Leu Lys Gln Leu  
 Press 2 to review your greeting and if you wish to re-record your greeting press 76 which will  
 delete your existing greeting, then press 5 to re-record. Press the # key to stop recording.  
 Ser Asn Gln Leu Arg Ser Asp Ile Ile Phe Gln Val Ser Arg Thr Gly  
 100 105 110  
 Gly His Leu Gly Ser Ser Leu Gly Val Val Gln Leu Thr Val Ala Leu  
 115 125  
 His Tyr Val Phe Asp Ala Pro Gln Asp Lys Ile Leu Trp Asp Val Gly  
 130 135 140  
 His Gln Tested on Tuesday 10 September and Sunday 10 October 1999 Asp Lys Met  
 145 150 155 160  
 Pro Thr Leu Arg Thr Met Asn Gly Leu Ser Phe Thr Lys Arg Ser  
 No Building Access, Building Access no Utilities, Building Access no IT, Building  
 Gln Ser Gln Tyr Asp Lys Phe Gly Ala Gly His Ser Ser Thr Ser Ile  
 Access no IPOL. Staff rang in to 568 0744 to receive message of availability. Calls  
 logged on voice mail.  
 Ser Ala Gly Leu Gly Met Ala Val Gly Arg Asp Leu Lys Gly Gln Asn  
 195 200 205  
 Asn His Val Ile Ser Val Ile Gly Asp Gly Ala Met Thr Ala Gly Gln  
 210 215 220  
 Ala Phe Gln Tested on Monday 4 October 1999  
 225 No IPOL available. All date stamping done manually, no problems. 240  
 Val Ile Leu Asn Asp Asn Lys Gln Val Ser Leu Pro Thr Ala Asn Leu  
 245 250 255  
 Asp Gly Pro Ile Pro Pro Val Gly Ala Leu Ser Ser Ala Leu Ser Lys  
 260 265 270  
 Leu Gln Ser Ser Lys Pro Leu Arg Gln Leu Arg Gln Val Ala Lys Gly  
 275 Entrance and Rear Courtyard door at 12.01am on 1 January 2000. Also instructions for  
 Val Thr Lys Gln Leu Gly Ala Pro Met His Gln Leu Ala Ala Lys Val  
 290 295 300  
 Asp Gln Tyr Ala Arg Gly Met Ile Ser Gly Ser Arg Ser Thr Leu Phe  
 305 310 315 320  
 Gln Gln Leu

<210> 193  
 <211> 88  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 193  
 Gly Gly His Leu Ser Ala Ser Leu Gly Val Val Gln Leu Thr Val Ala  
 1 5 10 15  
 Leu His Asn Val Phe Asn Ala Pro Gln Asp Lys Ile Val Trp Asp Val  
 20 25 30  
 Gly His Gln Thr Tyr Pro His Lys Ile Leu Thr Gly Arg Thr Arg  
 35 40 45  
 Met His Thr Ile Arg Lys Thr Ser Gly Leu Ala Gly Phe Pro Lys Arg  
 50 55 60  
 Asp Gln Ser Val Tyr Asp Thr Phe Gly Val Gly His Ser Ser Thr Ser  
 65 70 75 80  
 Ile Ser Ala Gly Leu Gly Met Ala  
 85

<210> 194

<211> 97  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 194  
 Pro Val Arg Glu Lys Leu Val Lys Ala Trp Arg Asn Asp Ser Glu Ile  
 1 5 10 15  
 Phe Ala His Tyr Gly Arg Leu Thr Thr Pro Tyr Ser Asp Glu Leu Leu  
 20 25 30  
 Gly Ser Lys Phe Cys Leu His Val Lys Gly Phe Glu Val Asn Thr Ala  
 35 40 45  
 Arg Ile Ala Asp Ser Leu Tyr Tyr Gly Cys Val Pro Val Ile Ile Ala  
 50 55 60  
 Asn His Tyr Asp Leu Pro Phe Ala Asp Ile Leu Asn Trp Lys Ser Phe  
 65 70 75 80  
 Ser Val Val Val Ala Thr Leu Asp Ile Pro Leu Leu Lys Arg Ile Leu  
 85 90 95  
 Lys

<210> 195  
 <211> 149  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 195  
 Gly Met His Thr Ser Lys Phe Cys Leu Asn Pro Ala Gly Asp Thr Pro  
 1 5 10 15  
 Ser Ala Cys Arg Leu Phe Asp Ala Ile Val Ser Leu Cys Ile Pro Val  
 20 25 30  
 Ile Val Ser Asp Ser Ile Glu Leu Pro Phe Glu Asp Val Ile Asp Tyr  
 35 40 45  
 Arg Lys Ile Ala Ile Phe Val Asp Thr Ala Thr Ser Leu Lys Arg Gly  
 50 55 60  
 Phe Leu Val Lys Leu Leu Arg Lys Val Arg Thr Glu Lys Ile Leu Glu  
 65 70 75 80  
 Tyr Gln Lys Glu Leu Lys Glu Val Lys Arg Phe Phe Glu Tyr Gly Asp  
 85 90 95  
 Pro Asn Gly Thr Val Lys Glu Ile Trp Arg Gln Ile Ser Gln Lys Leu  
 100 105 110  
 Pro Leu Ile Lys Leu Met Ile Asn Arg Asp Lys Arg Ile Val Lys Arg  
 115 120 125  
 Asp Met Ser Glu Pro Asp Cys Ser Cys Ile Cys Ser Asn Gln Thr Gly  
 130 135 140  
 Val Ile Ser Thr Leu  
 145

<210> 196  
 <211> 196  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 196  
 Met Ser Gln Val Ser Ala Thr Pro Cys Ala Pro Pro Asn Lys Glu Thr  
 1 5 10 15  
 Gly His Val Ile Glu Arg Arg Ser Ala Gly Tyr His Pro Ser Val Trp  
 20 25 30  
 Gly Asp Tyr Phe Leu Lys Tyr Asp Ser Pro Ser Asn Ser Val Lys Phe  
 35 40 45  
 Lys Phe Leu Gly Arg Val Glu Gly Gln Ile Glu Glu Leu Lys Gly Glu  
 50 55 60

Val Lys Lys Met Leu Ile Asp Val Val Asp Lys Pro Leu Pro Lys Leu  
 65 70 75 80  
 His Leu Ile Asp Gln Ile Gln Arg Leu Gly Ile Glu Tyr His Phe Glu  
 85 90 95  
 Arg Glu Val Asp Glu Gln Leu Glu Gln Ile His Lys Ser Tyr Ser Arg  
 100 105 110  
 Leu Asp His Glu Asp Phe Lys Val Asp Asp Leu His Thr Val Ala Leu  
 115 120 125  
 Ile Phe Arg Leu Leu Arg Gln His Gly Tyr Asn Ile Ser Ser Glu Ile  
 130 135 140  
 Phe Asp Lys Phe Lys Asp Ser Asn Gly Asn Phe Arg Glu Ser Leu Ile  
 145 150 155 160  
 Ser Asp Val Arg Gly Leu Leu Ser Leu Tyr Glu Ala Cys His Leu Arg  
 165 170 175  
 Cys His Gly Asp Ser Ile Leu Asp Glu Ala Leu Pro Phe Ala Thr Thr  
 180 185 190  
 His Leu Glu Ser  
 195

<210> 197  
 <211> 116  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 197  
 Met Ser Gln Val Ser Ala Thr Pro Cys Ala Pro Ser Asn Lys Gly Thr  
 1 5 10 15  
 Gly His Val Ile Glu Arg Arg Ser Ala Gly Tyr His Pro Ser Val Trp  
 20 25 30  
 Gly Asp Tyr Phe Leu Lys Tyr Asp Ser Pro Ser Asn Ser Val Lys Phe  
 35 40 45  
 Lys Phe Leu Gly Arg Val Glu Gly Gln Ile Glu Glu Leu Lys Gly Glu  
 50 55 60  
 Val Lys Lys Met Leu Thr Asp Ile Met Asp Lys Pro Leu Gln Lys Leu  
 65 70 75 80  
 His Leu Ile Asp Gln Ile Gln Arg Leu Gly Ile Glu Tyr His Phe Glu  
 85 90 95  
 Arg Glu Ile Asp Glu Gln Leu Glu Gln Ile His Lys Ser Tyr Ser Arg  
 100 105 110  
 Leu Asp His Glu  
 115

<210> 198  
 <211> 102  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 198  
 Met Ser Leu Pro Ile Ser Arg Val Pro Ser Ser Ser Pro Ala Glu Lys  
 1 5 10 15  
 Thr Ser Leu Val Pro Glu Gly Gly Ser Ala Ile Phe His Pro Thr Ile  
 20 25 30  
 Trp Ala Asp Tyr Phe Leu Lys His Ala Ser Asn Ser Asn Ser Thr Ser  
 35 40 45  
 Ser Asp Gly Val Val Glu Glu His Ile Glu Arg Leu Lys Gly Glu Val  
 50 55 60  
 Arg Lys Met Leu Met Gly Ala Met Asp Lys Pro Ser Gln Lys Leu Asn  
 65 70 75 80  
 Leu Ile Asp Gln Ile Gln Arg Leu Gly Phe Ala Tyr His Phe Glu His  
 85 90 95  
 Glu Ile Asp Glu Gln Leu

100

<210> 199  
 <211> 169  
 <212> PRT  
 <213> Eucalyptus grandis

&lt;400&gt; 199

```

Thr Ser Phe Leu Pro Ser Ser Ile His His Asn Gln Pro Ser Leu Leu
 1          5          10          15
Phe Phe Arg His Leu Cys Ser Ser Ser Ala Ala Thr Ser Ser Thr
          20          25          30
Ser Ser Gly Ala Gln Phe Val Thr Cys Thr Leu Lys Ile Glu Ala Gln
          35          40          45
Glu Ile Gly Arg Arg Ser Ala Asn Trp Gln Pro Asn Val Phe Asp Tyr
          50          55          60
Asp Phe Leu Gln Ser Leu Asn Val Asp Tyr Thr Glu Asp Lys Tyr Ser
          65          70          75          80
Glu Glu Ala Gln Arg Leu Lys Lys Glu Val Lys Gly Leu Phe Asp Lys
          85          90          95
Lys Met Asn Ser Val Ala Lys Leu Glu Phe Ile Asp Val Val Gln Arg
          100          105          110
Leu Gly Leu Gly Tyr Gln Phe Glu Thr Glu Ile Lys Asn Ala Leu Ser
          115          120          125
Ser Ile Tyr Asn Asn Ala Glu Asp Ala Gln Leu Leu Asp Asp Leu Tyr
          130          135          140
Ala Val Ser Leu Arg Phe Arg Leu Leu Arg Gln His Gly Phe Asn Ile
          145          150          155          160
Ser Gln Asp Ala Phe Gln Arg Phe Met
          165

```

<210> 200  
 <211> 132  
 <212> PRT  
 <213> Eucalyptus grandis

&lt;400&gt; 200

```

Ser Ile Arg Pro Asn Gln Pro Ser Leu Ser Leu Phe Ser Arg Pro Arg
 1          5          10          15
Ser Ser Phe Ser Ser Pro Ser Ala Val Ser Ser Gly Thr Arg Phe Ala
          20          25          30
Lys Cys Ala Leu Thr Ile Glu Asp Glu Asp Thr Ala Arg Arg Ser Ala
          35          40          45
Asn Trp Lys Pro Ser Val Trp Asp Tyr Gly Phe Val Gln Ser Leu Asn
          50          55          60
Thr Asp Phe Pro Val Asp Lys Tyr Thr Glu Gln Val Gln Arg Leu Lys
          65          70          75          80
Glu Glu Val Lys Gly Leu Phe His Arg Glu Met Asn Gln Val Ala Lys
          85          90          95
Leu Glu Phe Ile Asp Val Val Gln Arg Leu Gly Leu Gly Tyr His Phe
          100          105          110
Glu Thr Glu Ile Asn Asn Ser Leu Ser Ser Ile Tyr Asn Asn Thr Glu
          115          120          125
Asp Val Gln Leu
          130

```

<210> 201  
 <211> 116  
 <212> PRT  
 <213> Pinus radiata

&lt;400&gt; 201

```

Met Ala Ser Val Ser Val Lys Ala Gly Ala Thr Ser Thr Val Ser Cys
 1           5           10           15
Gly Leu Ala Ser Asn Asn Leu Ile Arg Arg Thr Ala Asn Pro His Pro
          20           25           30
Asn Val Trp Asp Tyr Asp Phe Val His Ser Leu Lys Ser Pro Tyr Asn
          35           40           45
Asp Ser Ser Tyr Thr Glu Arg Ala Glu Thr Leu Ile Gly Gln Leu Lys
          50           55           60
Val Met Leu Ser Ala Ala Ile Gly Gly Gly Glu Ser Met Ile Thr Pro
          65           70           75           80
Ser Ala Tyr Asp Thr Ala Trp Val Ala Arg Val Pro Ser Ile Asp Gly
          85           90           95
Ser Ala Cys Pro Gln Phe Pro Gln Thr Val Glu Trp Ile Leu Lys Asn
          100          105          110
Gln Leu Lys Asp
          115

```

&lt;210&gt; 202

&lt;211&gt; 121

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 202

```

Ala Ile Leu Ser Tyr Pro Pro Glu Ile Leu Ala Leu Pro Ser Pro Ser
 1           5           10           15
Phe Leu Tyr Ile Ser Ser Leu Ile Pro Met Ala Ser Val Val Asp Gln
          20           25           30
Ala Glu Leu Cys Ser Lys Ser Val Ser Met Ser Ser Pro Gly Val Gln
          35           40           45
Arg Arg Thr Gly Asp Tyr His Ser Asn Leu Trp Asp Asp Glu Phe Ile
          50           55           60
Gln Ser Leu Ser Thr Pro Tyr Gly Ala Pro Ser Tyr Arg Glu Arg Ala
          65           70           75           80
Asp Arg Leu Val Gly Glu Val Lys Glu Met Phe Asn Ser Leu Thr Val
          85           90           95
Leu Thr Pro His Asn Asp Leu Leu Glu Gln Leu Trp Met Val Asp Ser
          100          105          110
Val Glu Arg Leu Gly Ile Asp Arg His
          115          120

```

&lt;210&gt; 203

&lt;211&gt; 259

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 203

```

Asn Ile Gly Pro Ser Phe Leu Ser Ile Ser Ser Leu Val Arg Met Ala
 1           5           10           15
Ser Val Val Asp Gln Ala Glu Leu Cys Ser Lys Ser Val Ser Met Ser
          20           25           30
Ser Pro Gly Val Gln Arg Arg Thr Gly Asp Tyr His Ser Asn Leu Trp
          35           40           45
Asp Asp Asp Phe Ile Gln Ser Leu Ser Thr Pro Tyr Gly Ala Pro Ser
          50           55           60
Tyr Arg Glu Arg Ala Asp Arg Leu Val Gly Glu Val Lys Glu Met Phe
          65           70           75           80
Asn Ser Leu Thr Leu Leu Thr Pro Leu Asn Asp Leu Leu Gln Arg Leu
          85           90           95
Trp Met Val Asp Thr Val Glu Arg Leu Glu Ile Asp Arg His Phe Arg
          100          105          110

```

Asn Glu Ile Lys Ser Ala Leu Asp Tyr Val Tyr Ser Tyr Trp Ser Glu  
           115                          120                          125  
 Lys Gly Ile Gly Cys Gly Arg Glu Ser Val Val Thr Asp Leu Asn Ser  
           130                          135                          140  
 Thr Ala Leu Gly Phe Arg Thr Leu Arg Leu His Gly Phe Pro Val Ser  
           145                          150                          155                          160  
 Ser Asp Val Leu Glu Val Phe Lys Asp Gln Asn Gly Lys Phe Ala Gly  
                           165                          170                          175  
 Cys Ser Ala Asn Ala Glu Thr Glu Ala Glu Met Arg Asp Ile Leu Asn  
                           180                          185                          190  
 Leu Phe Arg Ala Ser Leu Val Ala Phe Pro Gly Glu Lys Val Met Glu  
           195                          200                          205  
 Glu Ala Gln Thr Phe Cys Thr Ser Tyr Leu Gln Glu Ala Leu Lys Thr  
           210                          215                          220  
 Val Pro Ile Ser Asn Asp Ser Leu Ser Arg Glu Ile Glu Tyr Val Ile  
           225                          230                          235                          240  
 Glu Tyr Gly Trp Leu Thr Asn Phe Ser Glu Ile Gly Ser Lys Glu Leu  
                           245                          250                          255  
 His Arg Arg

<210> 204  
 <211> 344  
 <212> PRT  
 <213> Pinus radiata

<400> 204  
 Ile Asp Val Phe Gly Glu Asp Thr Thr Phe Glu Thr Pro Tyr Leu Ile  
   1                          5                          10                          15  
 Arg Glu Lys Leu Leu Glu Leu Ala Lys Leu Glu Phe Asn Ile Phe His  
           20                          25                          30  
 Ser Leu Val Lys Arg Glu Leu Gln Ser Leu Leu Arg Trp Trp Lys Asp  
           35                          40                          45  
 Tyr Gly Phe Pro Glu Ile Thr Phe Ser Arg His Arg His Val Glu Tyr  
           50                          55                          60  
 Tyr Thr Leu Ala Ala Cys Ile Ala Asn Asp Pro Lys His Ser Ala Phe  
           65                          70                          75                          80  
 Arg Leu Gly Phe Gly Lys Ile Ser His Met Ile Thr Ile Leu Asp Asp  
                           85                          90                          95  
 Ile Tyr Asp Thr Phe Gly Thr Met Glu Glu Leu Glu Leu Leu Thr Ala  
           100                          105                          110  
 Ala Phe Lys Arg Trp Asp Pro Ser Ser Ile Glu Cys Leu Pro Asp Tyr  
           115                          120                          125  
 Met Lys Gly Val Tyr Met Ala Val Tyr Asp Asn Ile Asn Glu Met Ala  
           130                          135                          140  
 Arg Glu Ala Gln Lys Ile Gln Gly Trp Asp Thr Val Ser Tyr Ala Arg  
           145                          150                          155                          160  
 Lys Ser Trp Glu Ala Phe Ile Gly Ala Tyr Ile Gln Glu Ala Lys Trp  
                           165                          170                          175  
 Ile Ser Ser Gly Tyr Leu Pro Thr Phe Asp Glu Tyr Leu Glu Asn Gly  
           180                          185                          190  
 Lys Val Ser Phe Gly Ser Arg Ile Thr Thr Leu Glu Pro Met Leu Thr  
           195                          200                          205  
 Leu Gly Phe Pro Leu Pro Pro Arg Ile Leu Gln Glu Ile Asp Phe Pro  
           210                          215                          220  
 Pro Lys Phe Asn Asp Leu Ile Cys Ala Ile Leu Arg Leu Lys Gly Asp  
           225                          230                          235                          240  
 Thr Gln Cys Tyr Lys Ala Asp Arg Ala Arg Gly Glu Glu Ala Ser Ala  
                           245                          250                          255  
 Val Ser Cys Tyr Met Lys Asp His Pro Gly Ile Thr Glu Glu Asp Ala  
           260                          265                          270

Val Asn Gln Val Asn Ala Met Val Asp Asn Leu Thr Lys Glu Leu Asn  
 275 280 285  
 Trp Glu Leu Leu Arg Pro Asp Ser Gly Val Pro Ile Ser Tyr Lys Lys  
 290 295 300  
 Val Ala Phe Asp Ile Cys Arg Val Phe His Tyr Gly Tyr Lys Tyr Arg  
 305 310 315 320  
 Asp Gly Phe Ser Val Ala Ser Ile Glu Ile Lys Asn Leu Val Thr Arg  
 325 330 335  
 Thr Val Val Glu Thr Val Pro Leu  
 340

<210> 205  
 <211> 462  
 <212> PRT  
 <213> Pinus radiata

<400> 205  
 Arg Asp Ser Ala Phe Thr Asp Leu Asn Thr Thr Ala Leu Gly Phe Arg  
 1 5 10 15  
 Ile Phe Arg Leu His Gly Tyr Thr Val Ser Ser Asp Ala Phe Glu His  
 20 25 30  
 Phe Lys Asp Gln Met Gly Gln Phe Ser Ala Ser Ala Asn Asp Thr Glu  
 35 40 45  
 Leu Gln Ile Arg Ser Val Phe Asn Leu Phe Arg Ala Ser Leu Ile Ala  
 50 55 60  
 Phe Pro Glu Glu Lys Val Leu Glu Glu Ala Glu Asn Phe Ala Ala Ala  
 65 70 75 80  
 Tyr Leu Lys Ala Ala Leu Gln Thr Leu Pro Val Ser Gly Leu Ser Arg  
 85 90 95  
 Glu Ile Gln Tyr Val Phe Asp Tyr Arg Trp His Ser Asn Leu Pro Arg  
 100 105 110  
 Leu Glu Ala Arg Ser Tyr Val Asp Ile Leu Ala Asp Asn Thr Ile Ser  
 115 120 125  
 Gly Thr Pro Asp Ala Asn Thr Lys Lys Leu Leu Glu Leu Ala Lys Leu  
 130 135 140  
 Glu Phe Asn Ile Phe His Ser Leu Gln Gln Lys Glu Leu Gln Cys Leu  
 145 150 155 160  
 Trp Arg Trp Trp Lys Glu Trp Gly Cys Pro Glu Leu Thr Phe Val Arg  
 165 170 175  
 His Arg Tyr Val Glu Phe Tyr Thr Leu Val Ser Gly Thr Asp Met Val  
 180 185 190  
 Pro Glu His Ala Ala Phe Arg Leu Ser Phe Val Lys Thr Cys His Leu  
 195 200 205  
 Ile Thr Ile Leu Asp Asp Met Tyr Asp Thr Phe Gly Thr Ile Asp Glu  
 210 215 220  
 Leu Arg Leu Phe Thr Ala Ala Val Lys Arg Trp Asp Pro Ser Ala Thr  
 225 230 235 240  
 Glu Cys Leu Pro Glu Tyr Met Lys Gly Val Tyr Arg Val Leu Tyr Glu  
 245 250 255  
 Thr Val Asn Glu Met Ala Lys Glu Ala Gln Lys Ser Gln Gly Arg Asp  
 260 265 270  
 Thr Leu Gly Tyr Val Arg Gln Ala Leu Glu Asp Tyr Ile Gly Ser Tyr  
 275 280 285  
 Leu Lys Glu Ala Glu Trp Ile Ala Thr Gly Tyr Val Pro Thr Phe Gln  
 290 295 300  
 Glu Tyr Phe Glu Asn Gly Lys Leu Ser Ser Gly His Arg Ile Ala Thr  
 305 310 315 320  
 Leu Gln Pro Ile Leu Thr Leu Ser Ile Pro Phe Pro His His Ile Leu  
 325 330 335  
 Gln Glu Ile Asp Phe Pro Ser Lys Phe Asn Asp Tyr Ala Cys Ser Ile  
 340 345 350



Leu Arg Leu Arg Gly Asp Thr Arg Cys Tyr Lys Ala Asp Ser Ala Arg  
           355                                  360                  365  
 Gly Glu Glu Ala Ser Cys Ile Ser Cys Tyr Met Lys Glu Asn Pro Gly  
           370                                  375                  380  
 Ser Thr Gln Glu Asp Ala Leu His His Ile Asn Gly Met Ile Glu Asp  
 385                                  390                  395                  400  
 Met Ile Lys Lys Leu Asn Trp Glu Phe Leu Lys Pro Asp Asn Asn Ala  
                                   405                                  410                  415  
 Pro Ile Ser Ser Lys Lys Asn Ala Phe Asn Ile Ser Arg Gly Leu His  
                                   420                                  425                  430  
 His Phe Tyr Asn Tyr Arg Asp Gly Tyr Ser Val Ala Ser Asn Glu Thr  
                                   435                                  440                  445  
 Lys Asp Leu Val Ile Lys Thr Val Leu Glu Pro Val Leu Met  
           450                                  455                  460

&lt;210&gt; 206

&lt;211&gt; 100

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 206

Gly Ser Gln Leu Trp Asp Thr Ala Phe Ala Thr Gln Ala Ile Ile Ser  
 1                                  5                                  10                  15  
 Thr Asn Leu Ile Glu Glu Phe Gly Ser Thr Leu Gln Lys Ala His Thr  
                                   20                                  25                  30  
 Tyr Ile Lys Asn Ser Gln Val Leu Glu Asp Cys Pro Gly Asp Leu Asn  
                                   35                                  40                  45  
 Phe Trp Tyr Arg His Ile Ser Lys Gly Ala Trp Pro Phe Ser Thr Ala  
                                   50                                  55                  60  
 Asp His Gly Trp Pro Ile Ser Asp Cys Thr Ala Glu Gly Leu Lys Ala  
 65                                  70                                  75                  80  
 Ala Leu Val Leu Ser Lys Ile Pro Leu Glu Ile Val Gly Gln Pro Phe  
                                   85                                  90                  95  
 Arg Ser Tyr Gly  
                                   100

&lt;210&gt; 207

&lt;211&gt; 89

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 207

Met Trp Lys Leu Lys Val Ala Glu Gly Ala Asn Pro Trp Leu Arg Ser  
 1                                  5                                  10                  15  
 Leu Asn Asn His Val Gly Arg Gln Ile Trp Glu Phe Asp Pro Asn Cys  
                                   20                                  25                  30  
 Gly Ser Pro Glu Glu Ile Gln Glu Ile Glu Glu Ala Arg Ala Asn Phe  
                                   35                                  40                  45  
 Leu Lys His Arg Phe Glu Lys Lys His Ser Ser Asp Leu Met Met Arg  
                                   50                                  55                  60  
 Ile Gln Phe Ser Lys Glu Asn Thr Gly Arg Val Val Leu Pro Pro Val  
 65                                  70                                  75                  80  
 Lys Val Lys Asp Leu Asp Glu Ile Thr  
                                   85

&lt;210&gt; 208

&lt;211&gt; 198

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 208

Val Thr His Met Leu Arg Arg Ala Ile Ser Phe His Ser Thr Leu Gln  
 1 5 10 15  
 Ala His Asp Gly His Trp Pro Gly Asp Tyr Gly Gly Pro Met Phe Leu  
 20 25 30  
 Met Pro Gly Leu Val Ile Ala Leu Ser Ile Thr Gly Ala Leu Asn Ala  
 35 40 45  
 Val Leu Ser Glu Gln His Lys Gln Glu Met Cys Arg Tyr Leu Tyr Asn  
 50 55 60  
 His Gln Asn Lys Asp Gly Gly Trp Gly Leu His Ile Glu Gly Pro Ser  
 65 70 75 80  
 Thr Met Phe Gly Ser Val Leu Asn Tyr Val Thr Leu Arg Leu Leu Gly  
 85 90 95  
 Glu Ala Ala Asn Asp Gly Gln Gly Ala Met Glu Lys Ala Arg Lys Trp  
 100 105 110  
 Ile Leu Asp His Gly Ser Ala Thr Ala Ile Thr Ser Trp Gly Lys Met  
 115 120 125  
 Trp Leu Ser Val Leu Gly Ala Phe Glu Trp Ser Gly Asn Asn Pro Leu  
 130 135 140  
 Pro Pro Glu Ile Trp Leu Leu Pro Tyr Met Leu Pro Ile His Pro Gly  
 145 150 155 160  
 Arg Met Trp Cys His Cys Arg Met Val Tyr Leu Pro Met Ser Tyr Leu  
 165 170 175  
 Tyr Gly Lys Arg Phe Val Ser Pro Ile Thr Pro Thr Val Phe Val Leu  
 180 185 190  
 Glu Lys Arg Asn Phe Met  
 195

&lt;210&gt; 209

&lt;211&gt; 78

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 209

Met Trp Lys Leu Lys Ile Ala Glu Gly Gly Pro Trp Leu Thr Ser Val  
 1 5 10 15  
 Asn Asn His Val Gly Arg Gln His Trp Glu Phe Asp Pro Asp Ala Gly  
 20 25 30  
 Thr Pro Glu Glu Arg Ala Glu Val Glu Arg Val Arg Asp Glu Phe Thr  
 35 40 45  
 Arg Asn Arg Phe Arg Ile Lys Gln Ser Ala Asp Leu Leu Met Arg Met  
 50 55 60  
 Gln Leu Thr Lys Glu Asn Pro Ser Gly Pro Ile His Arg Arg  
 65 70 75

&lt;210&gt; 210

&lt;211&gt; 97

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 210

Tyr Val Trp Val Gly Glu Asp Gly Ile Lys Met Gln Ser Phe Gly Ser  
 1 5 10 15  
 Gln Ile Trp Asp Cys Gly Leu Ser Leu Gln Ala Leu Leu Ala Ser Asp  
 20 25 30  
 Leu Ile Asp Glu Ile Gly Pro Val Leu Lys Lys Gly His Glu Phe Leu  
 35 40 45  
 Lys Glu Ser Gln Ile Asp Arg Asn Pro Ser Gly Asp Leu Lys Lys Met  
 50 55 60  
 Tyr Arg His Ile Ser Lys Gly Ala Trp Ala Phe Ser Asp Lys Asp His  
 65 70 75 80  
 Gly Trp Gln Val Ser Asp Cys Thr Ala Glu Ser Met Lys Cys Cys Leu

Val 85 90 95

<210> 211  
 <211> 158  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 211

Met	Asp	Thr	Asp	Asn	Lys	Leu	Phe	Asn	Val	Gly	Val	Leu	Leu	Val	Ala
1				5					10					15	
Thr	Leu	Val	Val	Ala	Lys	Leu	Ile	Ser	Ala	Leu	Leu	Ile	Pro	Arg	Ser
			20					25					30		
Gly	Lys	Arg	Leu	Pro	Pro	Val	Val	Arg	Thr	Trp	Pro	Val	Val	Gly	Gly
		35				40					45				
Leu	Leu	Arg	Phe	Leu	Lys	Gly	Pro	Met	Val	Met	Leu	Arg	Glu	Glu	Tyr
	50					55				60					
Pro	Lys	Leu	Gly	Ser	Val	Phe	Thr	Leu	Asn	Leu	Leu	Asn	Lys	Lys	Ile
65					70					75					80
Thr	Phe	Phe	Ile	Gly	Pro	Glu	Val	Ser	Ala	His	Phe	Phe	Lys	Ala	Ser
			85					90					95		
Glu	Ser	Asp	Leu	Ser	Gln	Gln	Glu	Val	Tyr	Gln	Phe	Asn	Val	Pro	Thr
			100					105					110		
Phe	Gly	Pro	Gly	Val	Val	Phe	Asp	Val	Asp	Tyr	Thr	Ile	Arg	Gln	Glu
		115				120						125			
Gln	Phe	Arg	Phe	Phe	Thr	Glu	Ala	Leu	Arg	Ile	Asn	Lys	Leu	Lys	Gly
		130				135					140				
Tyr	Val	Asn	Gln	Met	Val	Met	Glu	Ala	Glu	Asp	Tyr	Phe	Ser		
145					150					155					

<210> 212  
 <211> 131  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 212

Met	Asp	Thr	Asp	Asn	Lys	Leu	Phe	Asn	Val	Gly	Val	Leu	Leu	Val	Ala
1				5					10					15	
Thr	Leu	Val	Val	Ala	Lys	Leu	Ile	Ser	Ala	Leu	Leu	Ile	Pro	Arg	Ser
			20					25					30		
Gly	Lys	Arg	Leu	Pro	Pro	Val	Val	Arg	Thr	Trp	Pro	Val	Val	Gly	Gly
		35				40					45				
Leu	Leu	Arg	Phe	Leu	Lys	Gly	Pro	Met	Val	Met	Leu	Arg	Glu	Glu	Tyr
	50					55				60					
Pro	Lys	Leu	Gly	Ser	Val	Phe	Thr	Leu	Asn	Leu	Leu	Asn	Lys	Lys	Ile
65					70					75					80
Thr	Phe	Phe	Ile	Gly	Pro	Glu	Val	Ser	Ala	His	Phe	Phe	Lys	Ala	Ser
			85					90					95		
Glu	Ser	Asp	Leu	Ser	Gln	Gln	Glu	Val	Tyr	Gln	Phe	Asn	Val	Pro	Thr
			100					105					110		
Phe	Gly	Pro	Gly	Val	Val	Phe	Asp	Val	Asp	Tyr	Thr	Ile	Arg	Gln	Glu
		115				120						125			
Gln	Phe	Arg													
130															

<210> 213  
 <211> 112  
 <212> PRT  
 <213> Eucalyptus grandis

&lt;400&gt; 213

Met Asp Thr Asp Asn Lys Leu Phe Asn Val Gly Val Leu Leu Val Ala  
 1 5 10 15  
 Thr Leu Val Val Ala Lys Leu Ile Ser Ala Ser Ile Pro Arg Ser Gly  
 20 25 30  
 Lys Arg Leu Pro Pro Val Val Arg Thr Trp Pro Val Val Gly Gly Leu  
 35 40 45  
 Leu Arg Phe Leu Lys Gly Pro Met Val Met Leu Arg Glu Glu Tyr Pro  
 50 55 60  
 Lys Leu Gly Ser Val Phe Thr Leu Asn Leu Leu Asn Lys Lys Ile Thr  
 65 70 75 80  
 Phe Phe Ile Gly Pro Glu Val Ser Ala His Phe Phe Lys Ala Ser Glu  
 85 90 95  
 Ser Asp Leu Ser Gln Gln Glu Val Tyr Gln Phe Asn Val Pro Thr Phe  
 100 105 110

&lt;210&gt; 214

&lt;211&gt; 152

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 214

Phe Leu Lys Gly Pro Met Val Met Leu Arg Glu Glu Tyr Pro Lys Leu  
 1 5 10 15  
 Gly Ser Val Phe Thr Leu Asn Leu Leu Asn Lys Lys Ile Thr Phe Phe  
 20 25 30  
 Ile Gly Pro Glu Val Ser Ala His Phe Phe Lys Ala Ser Glu Ser Asp  
 35 40 45  
 Leu Ser Gln Gln Glu Val Tyr Gln Phe Asn Val Pro Thr Phe Gly Pro  
 50 55 60  
 Gly Val Val Phe Asp Val Asp Tyr Thr Ile Arg Glu Glu Gln Phe Arg  
 65 70 75 80  
 Phe Phe Thr Glu Ala Leu Arg Ile Asn Lys Leu Lys Gly Tyr Val Asn  
 85 90 95  
 Gln Met Val Met Glu Ala Glu Asp Tyr Phe Ser Lys Trp Gly Asp Ser  
 100 105 110  
 Gly Glu Val Asp Leu Lys Tyr Glu Leu Glu His Leu Thr Ile Leu Thr  
 115 120 125  
 Ala Ser Arg Cys Leu Leu Gly Arg Glu Val Arg Glu Lys Leu Phe Asp  
 130 135 140  
 Asp Val Ser Ala Leu Phe His Asp  
 145 150

&lt;210&gt; 215

&lt;211&gt; 147

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 215

Phe Asp Asp Val Ser Ala Leu Phe His Asp Leu Asp Asn Gly Met Leu  
 1 5 10 15  
 Pro Ile Ser Val Ile Phe Pro Tyr Leu Pro Ile Pro Ala His His Arg  
 20 25 30  
 Arg Asp Lys Ala Arg Lys Lys Leu Ser Glu Ile Phe Ala Asn Ile Ile  
 35 40 45  
 Ser Ser Arg Lys Cys Ala Gly Lys Ser Glu Glu Asp Met Leu Gln Cys  
 50 55 60  
 Phe Ile Asp Ser Lys Tyr Lys Asn Gly Arg Pro Thr Thr Glu Ala Glu  
 65 70 75 80  
 Val Thr Gly Leu Leu Ile Ala Ala Leu Phe Ala Gly Gln His Thr Ser  
 85 90 95

Ser Ile Thr Ser Val Trp Thr Gly Ala Tyr Leu Leu Thr Asn Lys Lys  
 100 105 110  
 Tyr Leu Ser Ala Val Ser Asn Glu Gln Lys His Leu Met Glu Lys His  
 115 120 125  
 Gly Asn Asn Val Asp His Asp Val Leu Ser Glu Met Asp Val Leu Tyr  
 130 135 140  
 Arg Ser Ile  
 145

<210> 216  
 <211> 129  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 216  
 Tyr Leu Leu Thr Asn Lys Lys Tyr Leu Ser Ala Val Ser Asn Glu Gln  
 1 5 10 15  
 Lys His Leu Met Glu Lys His Gly Asn Asn Val Asp His Asp Val Leu  
 20 25 30  
 Ser Glu Met Asp Val Leu Tyr Arg Ser Ile Lys Glu Ala Leu Arg Leu  
 35 40 45  
 His Pro Pro Leu Ile Met Leu Leu Arg Ser Ser His Ser Asp Phe Ser  
 50 55 60  
 Val Lys Thr Arg Asp Gly Lys Glu Tyr Glu Val Gly Glu Val Ser Val  
 65 70 75 80  
 Leu Pro Trp Thr Leu Glu Ala Arg Lys Gly Val Gly Lys Ala Phe Ile  
 85 90 95  
 Thr Ala Phe Arg Ser Gly Ala Val Met Gly Phe Leu Leu Ala Ala Asn  
 100 105 110  
 Gly Leu Leu Val Leu Tyr Ile Ala Ile Asn Leu Phe Lys Ile Tyr Leu  
 115 120 125  
 Trp

<210> 217  
 <211> 118  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 217  
 Val Val Phe Asp Val Asp Tyr Thr Ile Arg Gln Glu Gln Phe Arg Phe  
 1 5 10 15  
 Phe Thr Glu Ala Leu Arg Ile Asn Lys Leu Lys Gly Tyr Val Asn Gln  
 20 25 30  
 Met Val Met Glu Ala Glu Asp Tyr Phe Ser Lys Trp Gly Asp Ser Gly  
 35 40 45  
 Glu Val Asp Leu Lys Tyr Glu Leu Glu His Leu Thr Ile Leu Thr Ala  
 50 55 60  
 Ser Arg Cys Leu Leu Gly Arg Glu Val Arg Glu Lys Leu Phe Asp Asp  
 65 70 75 80  
 Val Ser Ala Leu Phe His Asp Leu Asp Asn Gly Met Leu Pro Ile Ser  
 85 90 95  
 Val Ile Phe Pro Tyr Leu Pro Ile Pro Ala His His Arg Arg Asp Lys  
 100 105 110  
 Ala Arg Lys Lys Leu Ala  
 115

<210> 218  
 <211> 146  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 218  
 Ser Val Arg Arg Arg Ala Leu Glu Met Thr Thr Gly Arg Cys Leu Asp  
 1 5 10 15  
 Gly Leu Pro Leu Asp Gly Phe Asp Tyr Gly Ser Ile Leu Gly Gln Cys  
 20 25 30  
 Cys Glu Leu Leu Pro Ile Gly Tyr Val Gln Ile Pro Val Gly Val Ala Gly  
 35 40 45  
 Pro Leu Leu Leu Asp Gly Ile Glu Asn Met Val Pro Met Ala Thr Thr  
 50 55 60  
 Glu Gly Cys Leu Val Ala Ser Thr Asn Arg Gly Cys Lys Ala Ile His  
 65 70 75 80  
 Met Ser Gly Gly Ala Thr Ser Val Leu Leu Arg Asp Gly Met Thr Arg  
 85 90 95  
 Ala Pro Val Val Arg Phe Pro Thr Ala Arg Arg Ala Ala Gln Leu Lys  
 100 105 110  
 Phe Tyr Leu Glu Ala Pro Ile Thr Thr Lys Ala Cys Leu Ser Ser Ser  
 115 120 125  
 Thr Ala Pro Ser Lys Val Cys Gln Ala Cys Lys Gly Ile Gln Val Pro  
 130 135 140  
 Pro Ile  
 145

<210> 219  
 <211> 328  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 219  
 Val Ala Ser Tyr Ser Leu Glu Ser Ala Leu Gly Gly Asp Cys Arg Arg  
 1 5 10 15  
 Ala Ala Leu Val Arg Arg Arg Ala Leu Glu Ile Arg Thr Gly Arg Cys  
 20 25 30  
 Leu Asp Gly Leu Pro Leu Asp Gly Phe Asp Tyr Gly Ser Ile Leu Gly  
 35 40 45  
 Gln Cys Cys Glu Leu Pro Val Gly Tyr Val Gln Ile Pro Val Gly Val  
 50 55 60  
 Val Gly Pro Leu Leu Leu Asp Gly Leu Glu Asn Met Val Pro Met Ala  
 65 70 75 80  
 Thr Thr Glu Gly Cys Leu Val Ala Ser Ala Asn Arg Gly Cys Lys Ala  
 85 90 95  
 Ile His Met Ser Gly Gly Ala Thr Ser Val Leu Leu Arg Asp Gly Met  
 100 105 110  
 Thr Arg Ala Pro Val Val Arg Phe Pro Thr Ala Glu Arg Ala Ala His  
 115 120 125  
 Leu Lys Ser Tyr Leu Glu His Pro Lys Asn Phe Asp Ser Leu Ser Leu  
 130 135 140  
 Ile Phe Asn Ser Thr Ser Arg Phe Ala Arg Leu Gln Thr Ile Lys Cys  
 145 150 155 160  
 Ala Ile Ala Gly Arg Asn Leu Tyr Ile Arg Phe Ser Cys Phe Thr Gly  
 165 170 175  
 Asp Ala Met Gly Met Asn Met Val Ser Lys Gly Val Gln Asn Val Leu  
 180 185 190  
 Asp Phe Leu Gln Asn Glu Asn Pro Asp Met Asp Val Ile Ala Val Ser  
 195 200 205  
 Gly Asn Phe Cys Ala Asp Lys Lys Pro Thr Ala Val Asn Trp Ile Glu  
 210 215 220  
 Gly Arg Gly Lys Ser Val Val Cys Glu Ala Ile Ile Thr Glu Ala Val  
 225 230 235 240  
 Val Ser Lys Val Leu Lys Thr Thr Val Pro Ala Leu Leu Glu Leu Asn  
 245 250 255

Met Leu Lys Asn Leu Thr Gly Ser Ala Leu Ala Gly Ala Met Gly Gly  
                   260                  265                  270  
 Phe Asn Ala His Ala Ser Asn Ile Val Ser Ala Val Phe Ile Ala Thr  
                   275                  280                  285  
 Gly Gln Asp Pro Ala Gln Asn Ile Glu Ser Ser His Cys Ile Thr Met  
                   290                  295                  300  
 Met Glu Ala Ser Asn Asp Gly Lys Asp Leu His Val Ser Val Thr Met  
 305                  310                  315                  320  
 Pro Cys Ile Glu Val Gly Asn Ser  
                   325

<210> 220  
 <211> 175  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 220  
 Leu Gly Gly Asp Cys Arg Arg Ala Ala Ser Val Arg Arg Arg Ala Leu  
   1                  5                  10                  15  
 Glu Met Thr Thr Gly Arg Cys Leu Asp Gly Leu Pro Leu Asp Gly Phe  
                   20                  25                  30  
 Asp Tyr Gly Ser Ile Leu Gly Gln Cys Cys Glu Leu Pro Val Gly Tyr  
                   35                  40                  45  
 Val Gln Ile Pro Val Gly Val Ala Gly Pro Leu Leu Asp Gly Phe  
   50                  55                  60  
 Glu Ile Met Val Pro Met Ala Thr Thr Glu Gly Cys Leu Val Ala Ser  
 65                  70                  75                  80  
 Thr Asn Arg Gly Cys Lys Ala Ile His Met Ser Gly Gly Ala Thr Ser  
                   85                  90                  95  
 Val Leu Leu Arg Asp Gly Met Thr Arg Ala Pro Val Val Arg Phe Ser  
                   100                  105                  110  
 Thr Ala Arg Arg Ala Ala Gln Leu Lys Phe Tyr Leu Glu His Pro Asn  
                   115                  120                  125  
 Asn Tyr Lys Ser Leu Ser Leu Ile Phe Asn Ser Thr Ser Arg Phe Ala  
                   130                  135                  140  
 Arg Leu Gln Gly Ile Lys Cys Ala Ile Ala Gly Arg Asn Leu Tyr Met  
 145                  150                  155                  160  
 Arg Phe Cys Cys Ser Thr Gly Asp Ala Met Gly Asp Glu Tyr Gly  
                   165                  170                  175

<210> 221  
 <211> 220  
 <212> PRT  
 <213> Pinus radiata

<400> 221  
 Met Glu Ser Cys Gly Ser Gly Ile Ser Gly Thr Gly Lys Lys Met Lys  
   1                  5                  10                  15  
 Asn Ser Arg Thr Leu Ala Ser Asp Ala Leu Pro Leu Pro Val Gly Leu  
                   20                  25                  30  
 Thr Asn Lys Val Phe Phe Ile Leu Phe Phe Thr Ala Ser Tyr Phe Leu  
                   35                  40                  45  
 Met Arg Arg Trp Arg Glu Lys Ile Arg Thr Ser Thr Pro Leu His Val  
   50                  55                  60  
 Leu Ser Leu Gly Glu Leu Val Ala Ile Val Ala Gln Leu Ala Ser Phe  
 65                  70                  75                  80  
 Ile Tyr Leu Leu Gly Phe Phe Gly Ile Asp Tyr Val Gln Asn Phe Ile  
                   85                  90                  95  
 Thr Gly Gly Asn Asp Asp Asp Ala Arg Glu Asp Asp Lys Leu Arg  
                   100                  105                  110  
 Ser Pro Val Pro Lys Glu Ala Val Ala Ile Arg Pro Ser Ala Pro Gln

115	120	125
Val Gln Leu Asn Gly Ile Ser	Leu Gly Asp Asn Lys Asp Asp Asp Ile	
130	135	140
Ala Ala Ala Val Cys Asn Gly Thr Val Ala Ser Tyr Ser Leu Glu Ser		
145	150	155
Ser Leu Gly Asp Cys Met Arg Ser Ala Arg Val Arg Arg Arg Ser Leu		
165	170	175
Glu Met Met Thr Gly Arg Ser Leu Asp Gly Leu Pro Leu Glu Gly Phe		
180	185	190
Asp Tyr Gly Ser Ile Leu Gly Gln Cys Cys Glu Leu Pro Val Gly Tyr		
195	200	205
Val Gln Ile Pro Val Gly Val Ala Gly Pro Leu Leu		
210	215	220

<210> 222  
 <211> 91  
 <212> PRT  
 <213> Pinus radiata

<400> 222
Asp Leu His Ile Ser Val Thr Met Pro Cys Ile Glu Val Gly Thr Val
1 5 10 15
Gly Gly Gly Thr Gln Leu Ala Ser Gln Ser Ala Cys Leu Asn Leu Ile
20 25 30
Gly Val Lys Gly Ala Asn Val Gln Ser Pro Gly Ala Asn Ala Arg Leu
35 40 45
Leu Ala Arg Ile Val Ala Gly Ala Val Leu Ala Gly Glu Leu Ser Leu
50 55 60
Met Ser Ala Leu Ala Ala Gly Gln Leu Val Lys Ser His Met Lys Tyr
65 70 75 80
Asn Arg Ser Ile Lys Asp Ile Lys Ala Ile Ser
85 90

<210> 223  
 <211> 187  
 <212> PRT  
 <213> Pinus radiata

<400> 223
Ser Phe Glu Ile His Thr Gly Lys Ser Ala Asp Ile Ser Arg Ala Gln
1 5 10 15
Ser Ala Tyr Thr Gln Gln Asn Asn Asn Ile Phe Thr Ser Ser Lys Ile
20 25 30
His Pro Val Val Ile Val Pro Gly Thr Gly Gly Asn Gln Val Glu Ala
35 40 45
Arg Leu Thr Ala Asp Tyr Lys Pro Ser Gly Leu Leu Cys Arg Arg Trp
50 55 60
Asn Trp Glu Arg Glu Trp Phe Arg Ile Trp Phe Asp Val Pro Val Val
65 70 75 80
Leu Pro Pro Leu Thr Gln Cys Phe Ala Asp Arg Ile Ser Leu Val Tyr
85 90 95
Asp Pro His Thr Asp Glu Tyr Tyr Asn Ala Pro Gly Val Glu Thr Arg
100 105 110
Val Pro Tyr Phe Gly Ser Thr Glu Gly Met Lys Tyr Leu Asp Pro Cys
115 120 125
Phe Lys Tyr Ile Thr Pro Tyr Met Ser Ser Leu Val Lys Ser Leu Glu
130 135 140
Asp Val Gly Tyr Val Asp Gly Lys Ser Leu Phe Gly Ala Pro Tyr Asp
145 150 155 160
Phe Arg Tyr Gly Pro Gly Thr Lys Ser Ser Ser Val Gly Ala Lys Tyr
165 170 175



Leu Glu Asn Leu Arg Lys Leu Val Glu Glu Ala  
 180 185

<210> 224  
 <211> 117  
 <212> PRT  
 <213> Pinus radiata

<400> 224  
 Ser Ala Leu Ile Ile Gly Ser Phe Ile Phe Cys Ile Phe Leu Tyr Ile  
 1 5 10 15  
 Lys Gly His Val Ala Pro Ser Ser Thr Asp Ser Gly Ser Ser Gly Asn  
 20 25 30  
 Val Val Ile Asp Phe Tyr Trp Gly Met Glu Leu Tyr Pro Arg Ile Gly  
 35 40 45  
 Lys Asn Phe Asp Ile Lys Val Phe Thr Asn Cys Arg Phe Gly Met Met  
 50 55 60  
 Ser Trp Ala Val Leu Ala Val Thr Tyr Ser Ile Lys Gln Tyr Glu Glu  
 65 70 75 80  
 Tyr Gly Arg Val Ala Asp Ser Met Leu Val Ser Ser Ile Leu Met Val  
 85 90 95  
 Val Tyr Val Thr Lys Val Leu Leu Val Gly Ile Trp Leu Leu Glu His  
 100 105 110  
 His Gly Tyr Asn Ser  
 115

<210> 225  
 <211> 210  
 <212> PRT  
 <213> Pinus radiata

<400> 225  
 Phe Ala Val Val Gly Pro Leu Gln Leu Thr Ser Tyr Pro Leu Ile Lys  
 1 5 10 15  
 Leu Val Gly Ile Arg Thr Gly Leu Pro Leu Pro Ser Leu Trp Glu Ile  
 20 25 30  
 Phe Ala Gln Leu Ala Val Tyr Phe Met Val Glu Asp Tyr Gly Asn Tyr  
 35 40 45  
 Trp Ile His Arg Trp Leu His Cys Lys Trp Gly Tyr Glu Lys Ile His  
 50 55 60  
 His Val His His Glu Phe Thr Ala Pro Met Gly Phe Ala Ala Pro Tyr  
 65 70 75 80  
 Ala His Trp Ser Glu Val Leu Ile Leu Gly Ile Pro Thr Phe Val Gly  
 85 90 95  
 Pro Ala Ile Ala Pro Gly His Met Ile Thr Phe Trp Cys Trp Val Val  
 100 105 110  
 Leu Arg Gln Val Glu Ala Ile Glu Thr His Ser Gly Tyr Asp Phe Pro  
 115 120 125  
 Trp Thr Leu Thr Lys Leu Ile Pro Phe Tyr Gly Gly Ala Glu Tyr His  
 130 135 140  
 Asp Tyr His His Tyr Val Gly Gly Gln Ser Gln Ser Asn Phe Ala Ser  
 145 150 155 160  
 Val Phe Thr Tyr Cys Asp Tyr Leu Tyr Gly Thr Asp Lys Gly Tyr Arg  
 165 170 175  
 Tyr Arg Lys Glu His Leu Leu Lys Ala Arg Glu Phe Glu Tyr Arg Leu  
 180 185 190  
 Lys Gln Met Ile Leu Arg Lys Lys Thr Ala Met Glu Gln Phe Gln Ile  
 195 200 205  
 Ser Leu  
 210

<210> 226  
 <211> 86  
 <212> PRT  
 <213> Pinus radiata

<400> 226  
 Gly Pro His Leu Phe Thr Leu Trp Leu Trp Met Ser Leu Arg Val Leu  
 1 5 10 15  
 Glu Thr Val Glu Ala His Cys Gly Tyr Asp Phe Pro Trp Ser Ile Ser  
 20 25 30  
 Lys Leu Phe Pro Leu Tyr Gly Gly Ala Asp Phe His Asp Tyr His His  
 35 40 45  
 Arg Leu Leu Tyr Thr Lys Ser Gly Asn Tyr Ser Ser Thr Phe Thr Tyr  
 50 55 60  
 Met Asp Trp Leu Phe Gly Thr Asp Lys Gly Tyr Arg Lys Leu Lys Gly  
 65 70 75 80  
 Leu Gln Lys Asp Ser Lys  
 85

<210> 227  
 <211> 141  
 <212> PRT  
 <213> Pinus radiata

<400> 227  
 Met Ala Thr Leu Val Glu Arg Gly Trp Leu Tyr Leu Ile Thr Asn Phe  
 1 5 10 15  
 Thr Asp Phe Gln Leu Ala Ser Ile Gly Ser Phe Leu Leu His Glu Ser  
 20 25 30  
 Ile Phe Tyr Leu Ser Gly Leu Pro Phe Ile Leu Leu Glu Thr Thr Gly  
 35 40 45  
 Leu Leu Ser Lys Tyr Lys Ile Gln Ser Lys Thr Asn Thr Val Ala Ala  
 50 55 60  
 Gln Glu Lys Cys Ile Thr Arg Leu Leu Leu Tyr His Phe Cys Val Asn  
 65 70 75 80  
 Leu Pro Val Met Val Ser Tyr Pro Val Phe Arg Phe Met Gly Met  
 85 90 95  
 Thr Ser Val Leu Pro Leu Pro Ser Trp Lys Val Val Val Ser Gln Leu  
 100 105 110  
 Val Cys Tyr Phe Ile Leu Glu Asp Phe Val Phe Tyr Trp Gly His Arg  
 115 120 125  
 Ile Leu His Ser Lys Trp Leu Tyr Lys His Val His Ser  
 130 135 140

<210> 228  
 <211> 381  
 <212> PRT  
 <213> Pinus radiata

<400> 228  
 Met Gly Glu Glu Leu Gln Thr Trp Ile Leu Met Val Thr Ala Arg Ala  
 1 5 10 15  
 Pro Thr Asn Ile Ala Val Ile Lys Tyr Trp Gly Lys Arg Asp Glu Lys  
 20 25 30  
 Leu Ile Leu Pro Ile Asn Asp Ser Ile Ser Phe Thr Leu Asp Pro Asp  
 35 40 45  
 His Leu Ser Ala Thr Thr Thr Val Ala Val Ser Pro Ser Phe Thr Ser  
 50 55 60  
 Asp Arg Met Trp Leu Asn Gly Lys Glu Val Ser Leu Gly Gly Glu Arg  
 65 70 75 80  
 Tyr Gln Asn Cys Leu Arg Glu Ile Arg Ser Arg Gly Asn Asp Val Val

```
<210> 229
<211> 81
<212> PRT
<213> Pinus radiata
```

```
<210> 230
<211> 189
<212> PRT
<213> Pinus radiata
```

&lt;400&gt; 230

```

Met Pro Leu Thr Leu Leu Leu Ala Asn Thr Trp Ala Ser Ser Ala Ile
 1           5           10           15
Val Ser Arg Arg Val Ser Leu Phe Val Ala Cys Ser Thr Thr Val Val
      20           25           30
Ser Arg Ser Phe Ser Lys Ser Cys Ser Gly Ala Ile Pro Arg Lys Pro
      35           40           45
Lys Ser Ala His Pro Ala Leu Thr Gly Ser Arg Thr Cys Phe Ser Arg
      50           55           60
Asn Pro Ile Val Arg Asn Leu Ile Gly Ser Ala Ser Lys Met Gly Ala
65           70           75           80
Thr Val Glu Asp Thr Thr Met Asp Ala Val Gln Arg Arg Leu Met Phe
      85           90           95
Glu Asp Glu Cys Ile Leu Val Asp Glu Glu Asp His Val Ile Gly His
      100          105          110
Asp Ser Lys Tyr Asn Cys His Leu Met Glu Lys Ile Glu Ser Glu Asn
      115          120          125
Leu Leu His Arg Ala Phe Ser Val Phe Leu Phe Asn Thr Lys Tyr Glu
      130          135          140
Leu Leu Leu Gln Gln Arg Ser Ala Thr Lys Val Thr Phe Pro Leu Val
      145          150          155          160
Trp Thr Asn Thr Cys Cys Ser His Pro Leu Tyr Arg Glu Ser Glu Leu
      165          170          175
Ile Glu Glu Asn Asn Leu Gly Ser Glu Met Gln Pro Lys
      180          185

```

&lt;210&gt; 231

&lt;211&gt; 113

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 231

```

Met Ala Gly Ile Pro Val Leu Arg Pro Phe Cys Ile Cys Leu Leu Ser
 1           5           10           15
Val Tyr Met Leu His Ile Val Ala Ala Val Ala Ser Pro Arg Leu Gly
      20           25           30
Arg Ser Ser Phe Pro Arg Gly Phe Lys Phe Gly Ala Gly Ser Ser Ala
      35           40           45
Tyr Gln Ala Glu Gly Ala Ala His Glu Gly Gly Lys Gly Pro Ser Ile
      50           55           60
Trp Asp Thr Phe Ser His Thr Pro Gly Lys Ile Ala Asp Gly Lys Asn
65           70           75           80
Gly Asp Val Ala Val Asp Gln Tyr His Arg Tyr Lys Glu Asp Val Gln
      85           90           95
Leu Leu Lys Tyr Met Gly Met Asp Val Tyr Arg Phe Ser Ile Ser Trp
      100          105          110
Ser

```

&lt;210&gt; 232

&lt;211&gt; 127

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 232

```

Gly Pro Ser Ile Trp Asp Thr Phe Ser His Thr Pro Gly Lys Ile Ala
 1           5           10           15
Asp Gly Lys Asn Gly Asp Val Ala Val Asp Gln Tyr His Arg Tyr Lys
      20           25           30
Glu Asp Val Gln Leu Leu Lys Asn Met Gly Met Asp Val Tyr Arg Phe

```

```
<210> 233
<211> 118
<212> PRT
<213> Eucalyptus grandis
```

```
<210> 234
<211> 111
<212> PRT
<213> Pinus radiata
```

```
<210> 235
<211> 391
<212> PRT
<213> Pinus radiata
```

&lt;400&gt; 235

```

Met Met Gln Lys Tyr Ile Gly Ala Asp Val Thr Ser Met Val Thr Leu
 1           5           10           15
Pro Val Ile Ile Phe Glu Pro Met Thr Met Leu Gln Lys Ser Ala Glu
          20           25           30
Leu Met Glu Tyr Thr Tyr Leu Leu Asp Met Ala Asp Glu Cys Glu Asp
      35           40           45
Pro Tyr Leu Lys Met Ala Tyr Ala Ala Ser Trp Ala Ile Ser Val Tyr
      50           55           60
Pro Ala Tyr Gln Arg Ser Trp Lys Pro Phe Asn Pro Ile Leu Gly Glu
65           70           75           80
Thr Tyr Glu Met Val Asn His Gly Gly Ile Thr Phe Ile Ala Glu Gln
          85           90           95
Val Ser His His Pro Pro Met Gly Ser Ala Tyr Ala Glu Asn Glu His
          100          105          110
Phe Thr Tyr Ser Leu Ser Ser Lys Val Lys Thr Lys Phe Leu Gly Asn
          115          120          125
Ser Val Asp Ile Tyr Pro Leu Gly Arg Thr Arg Val Val Leu Lys Lys
          130          135          140
Ser Gly Asp Val Leu Asp Leu Val Pro Pro Pro Ser Lys Val His Asn
145          150          155          160
Leu Ile Phe Gly Arg Thr Trp Ile Asp Ser Pro Gly Glu Met Val Leu
          165          170          175
Thr Asn Leu Thr Thr Gly Asp Lys Val Val Leu Tyr Phe Gln Pro Cys
          180          185          190
Gly Trp Phe Gly Ala Gly Arg Tyr Glu Val Asp Gly Tyr Val Tyr Asp
          195          200          205
Ser Lys Glu Glu Pro Lys Ile Leu Met Thr Gly Lys Trp Asn Arg Ser
          210          215          220
Met Gly Tyr Gln Pro Cys Asp Ala Glu Gly Glu Pro Leu Pro Gly Thr
225          230          235          240
Glu Leu Lys Glu Val Trp Arg Val Ala Asp Leu Pro Lys Asn Asp Lys
          245          250          255
Phe Gln Tyr Thr Tyr Phe Ala His Lys Ile Asn Ser Phe Asp Thr Ala
          260          265          270
Pro Lys Lys Leu Leu Ala Ser Asp Ser Arg Leu Arg Pro Asp Arg Ser
          275          280          285
Ala Leu Glu Met Gly Asp Leu Ser Lys Ala Gly Ala Glu Lys Ser Asn
          290          295          300
Leu Glu Glu Arg Gln Arg Ala Glu Lys Arg Cys Arg Glu Glu Lys Asn
305          310          315          320
Gln Pro Phe Thr Pro Arg Trp Phe Thr Val Thr Gly Glu Val Ala Thr
          325          330          335
Thr Pro Trp Gly Asp Leu Glu Val Tyr Glu Tyr Asn Gly Lys Tyr Ser
          340          345          350
Glu His Arg Ala Ser Val Asp Asp Ser Asn Phe Asp Asp Gly Thr Asp
          355          360          365
Ser Lys Ser Met Glu Phe Asn Pro Trp Gln Tyr Gly Asn Ile Glu Ser
          370          375          380
Gly Ser Asn Lys Lys Val Glu
385          390

```

&lt;210&gt; 236

&lt;211&gt; 27

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 236

```

Met Met Gln Lys Tyr Ile Gly Ala Asp Val Thr Ser Met Val Thr Leu
 1           5           10           15
Pro Val Ile Ile Phe Glu Pro Met Thr Met Leu

```

20

25

<210> 237  
 <211> 134  
 <212> PRT  
 <213> Pinus radiata

&lt;400&gt; 237

Tyr Leu Val Leu Ile Ser Gln Leu Arg Val Gly Met Asp Leu Ser Lys  
 1 5 10 15  
 Val Thr Phe Pro Thr Phe Val Leu Glu Pro Arg Ser Met Leu Glu Arg  
 20 25 30  
 Ile Thr Asp Phe Met Ser His Pro Asp Leu Ile Phe Gly Ala Glu Asn  
 35 40 45  
 Ser Asn Asp Pro Glu Glu Arg Phe Met Arg Val Leu Ser Tyr Tyr Leu  
 50 55 60  
 Ala Gly Trp His Ile Lys Pro Lys Gly Val Lys Lys Pro Tyr Asn Pro  
 65 70 75 80  
 Val Leu Gly Glu Phe Phe Arg Cys Arg Tyr Asp Tyr Ser Asn Asn Thr  
 85 90 95  
 Gln Gly Phe Tyr Ile Ala Glu Gln Val Ser His His Pro Pro Ile Ser  
 100 105 110  
 Ala Phe Phe Tyr Ile Ser Pro Ala Asn Arg Val Ser Ile Ile Gly Glu  
 115 120 125  
 Leu Arg Pro Lys Ser Lys  
 130

<210> 238  
 <211> 133  
 <212> PRT  
 <213> Eucalyptus grandis

&lt;400&gt; 238

Ser Ser Lys Gly Arg His Cys Lys Pro Phe Asn Pro Leu Leu Gly Glu  
 1 5 10 15  
 Thr Tyr Glu Ala Asp Tyr Pro Glu Arg Gly Val His Phe Phe Ser Glu  
 20 25 30  
 Lys Val Ser His His Pro Thr Leu Ile Ala Cys His Cys Glu Gly Arg  
 35 40 45  
 Gly Trp Lys Phe Trp Ala Asp Ser Asn Leu Arg Thr Lys Phe Trp Gly  
 50 55 60  
 Gln Ser Ile Gln Leu Asp Pro Val Gly Ala Leu Thr Leu Glu Phe Asp  
 65 70 75 80  
 Asp Gly Glu Ile Phe Gln Trp Asn Lys Val Thr Thr Ser Ile Asn Asn  
 85 90 95  
 Leu Ile Ile Gly Lys Val Tyr Cys Asp His His Gly Val Met Asn Ile  
 100 105 110  
 His Gly Asn His Gln Tyr Ser Cys Lys Leu Lys Phe Lys Glu Pro Ser  
 115 120 125  
 Ile Leu Ala Glu Leu  
 130

<210> 239  
 <211> 116  
 <212> PRT  
 <213> Eucalyptus grandis

&lt;400&gt; 239

Arg Thr Cys Asp Trp Ser Met Arg Ala Ser Trp Ala Ile Ser Val Tyr  
 1 5 10 15  
 Tyr Ala Tyr Gln Arg Thr Trp Lys Pro Phe Asn Pro Ile Leu Gly Glu

20 25 30  
 Thr Tyr Glu Leu Ala Asn His Gly Gly Ile Thr Phe Ile Ala Glu Gln  
 35 40 45  
 Val Cys His His Pro Pro Met Ser Ala Gly His Ala Glu Asn Asp His  
 50 55 60  
 Phe Thr Tyr Asp Val Thr Ser Lys Leu Lys Thr Lys Phe Leu Gly Asn  
 65 70 75 80  
 Ser Val Asp Val Tyr Pro Val Gly Arg Thr Arg Val Thr Leu Lys Arg  
 85 90 95  
 Asp Gly Val Val Leu Asp Leu Val Pro Pro Thr Lys Val Asn Asn  
 100 105 110  
 Leu Ile Phe Gly  
 115

<210> 240  
 <211> 105  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 240  
 Ser Arg Leu Arg Pro Asp Arg Tyr Ala Leu Glu Pro Gly Asp Leu Pro  
 1 5 10 15  
 Lys Ala Gly Ala Glu Lys Ser Ser Leu Glu Glu Arg Gln Arg Gly Glu  
 20 25 30  
 Lys Lys Asn Arg Glu Met Lys Gly Gln Lys Phe Thr Pro Arg Trp Phe  
 35 40 45  
 Asp Leu Thr Asp Glu Ile Ser Pro Thr Pro Trp Gly Asp Leu Glu Val  
 50 55 60  
 Tyr Arg Tyr Asn Gly Lys Tyr Thr Glu His Arg Ala Val Val Asp Ser  
 65 70 75 80  
 Leu Asp Thr Ile Glu Glu Ser Asp Ile Gln Ser Thr Glu Phe Asn Pro  
 85 90 95  
 Trp Gln Tyr Glu Ala Thr Phe Ala Glu  
 100 105

<210> 241  
 <211> 117  
 <212> PRT  
 <213> Pinus radiata

<400> 241  
 Val Leu Arg Gly Leu Asp Thr Val Glu Asp Asp Thr Ser Ile Pro Leu  
 1 5 10 15  
 Asp Thr Lys Leu Pro Ile Leu Lys Ala Phe Tyr Lys His Ile Tyr Asp  
 20 25 30  
 Pro Ser Trp His Phe Ser Cys Gly Val Glu His Tyr Lys Glu Leu Met  
 35 40 45  
 Glu Lys Phe His His Val Ser Thr Thr Phe Leu Arg Leu Gly Arg Gly  
 50 55 60  
 Tyr Gln Glu Ala Ile Glu Glu Ile Thr Lys Lys Met Gly Ala Gly Met  
 65 70 75 80  
 Ala Lys Phe Ile Cys Lys Glu Val Glu Ser Val Glu Asp Tyr Asp Glu  
 85 90 95  
 Tyr Cys His Tyr Val Ala Gly Leu Val Gly Phe Gly Leu Ser Arg Leu  
 100 105 110  
 Phe His Ala Ala Gln  
 115

<210> 242  
 <211> 190  
 <212> PRT



&lt;213&gt; Pinus radiata

&lt;400&gt; 242

```

Met Ala Ile Tyr Thr Pro Gln Pro Ala His Arg Leu Ile Ser Trp Ser
 1          5          10          15
Thr Met Glu Asn His Thr Val Val Ile Ala Ala Ala Ile Ser Phe Val
 20          25          30
Ser Val Leu Leu Ser Tyr Tyr Ile Val Leu Ser Arg Trp Lys Arg Arg
 35          40          45
Ser Asn Gly Leu Arg Gly Ile Gln Ser Lys Ser Phe Glu Lys Ser Thr
 50          55          60
Asp Asp Asn Gly Ile Ala Ile Glu Ala Ala Gly Gly Thr Asp Val Ile
 65          70          75          80
Ile Val Gly Ala Gly Val Ala Gly Ser Ala Leu Ala Tyr Thr Leu Gly
 85          90          95
Lys Asp Gly Arg Arg Ile His Val Ile Glu Arg Asp Leu Ser Glu Pro
 100          105          110
Asp Arg Ile Val Gly Glu Leu Leu Gln Pro Gly Gly Tyr Leu Lys Leu
 115          120          125
Ile Glu Leu Gly Leu Gln Asp Cys Val Glu Gly Ile Asp Ala Gln Ser
 130          135          140
Ile Phe Gly Asp Ala Leu Phe Lys Glu Gly Lys Asp Thr Lys Val Ala
 145          150          155          160
Tyr Pro Leu Glu Asn His His Ala Asp Arg Ala Gly Arg Ser Phe His
 165          170          175
Asn Gly Arg Phe Ile Gln Arg Met Arg Glu Lys Ala Ala Ser
 180          185          190

```

&lt;210&gt; 243

&lt;211&gt; 124

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 243

```

Cys Leu Thr Thr Asp Ser Gly Gln Val Ile Asn Cys Arg Asn Arg Tyr
 1          5          10          15
Thr Ala Met Ala Ile Tyr Thr Pro Gln Pro Ala His Arg Leu Ile Ser
 20          25          30
Trp Ser Thr Met Glu Asn His Thr Val Ala Ile Ala Val Ala Ile Gly
 35          40          45
Phe Val Ser Val Leu Leu Ser Tyr Tyr Ile Val Leu Asn Arg Trp Lys
 50          55          60
Arg Arg Ser Asn Gly Leu Arg Gly Ile Gln Ser Lys Ser Phe Glu Lys
 65          70          75          80
Ser Thr Asp Asp Asn Gly Ile Ala Ile Glu Ala Ala Gly Gly Thr Asp
 85          90          95
Val Ile Ile Val Gly Ala Gly Val Ala Gly Ser Ala Leu Ala Tyr Thr
 100          105          110
Leu Gly Lys Asp Gly Arg Arg Ile His Val Ile Glu
 115          120

```

&lt;210&gt; 244

&lt;211&gt; 123

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 244

```

Met Asp Gly Gln Tyr Leu Val Ser Gly Val Leu Ala Leu Phe Leu Gly
 1          5          10          15
Ile Phe Leu Leu Tyr Lys Gly Leu Gly Lys Gln Lys Arg Arg Leu Ser
 20          25          30

```

Lys Lys Gly Arg Gly Asp Asp Tyr Val Lys Ser Ser Val Asp Gly Gly  
           35                          40                          45  
 Phe Val Pro Gly Gly Val Asp Gly Ser Thr Asp Ile Val Ile Val Gly  
           50                          55                          60  
 Ala Gly Val Ala Gly Ala Ala Leu Ala Tyr Ala Leu Gly Lys Asp Gly  
   65                          70                          75                          80  
 Arg Arg Val Arg Val Ile Glu Arg Asp Leu Thr Glu Gln Asp Arg Ile  
                           85                          90                          95  
 Val Gly Glu Leu Leu Gln Pro Gly Gly Tyr Leu Lys Leu Met Glu Leu  
                           100                          105                          110  
 Asp Leu Ala Asp Cys Val Gln Thr Ile Asp Ala  
           115                          120

<210> 245  
 <211> 221  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 245  
 Leu Gly Ser Lys Tyr Lys Pro Gln Glu Glu Phe Val Glu Trp Ile Gln  
   1                          5                          10                          15  
 Lys Gly Thr Lys Pro Ile Tyr Ile Gly Phe Gly Ser Met Pro Leu Glu  
           20                          25                          30  
 Asp Pro Lys Lys Thr Thr Asp Ile Ile Ile Lys Ala Leu Thr Asp Thr  
           35                          40                          45  
 Gly Gln Arg Gly Ile Val Gly Arg Gly Trp Gly Asp Leu Gly Thr Leu  
   50                          55                          60  
 Leu Asp Val Pro Asp Ser Val Phe Leu Leu Glu Asp Cys Pro His Asp  
   65                          70                          75                          80  
 Trp Leu Phe Pro Gln Cys Ser Ala Val Val His His Gly Gly Ala Gly  
                           85                          90                          95  
 Thr Thr Ala Thr Gly Leu Lys Ala Gly Cys Pro Thr Thr Ile Val Pro  
           100                          105                          110  
 Phe Phe Gly Asp Gln Phe Phe Trp Gly Asp Arg Val His Gln Arg Gly  
           115                          120                          125  
 Leu Gly Pro Ala Pro Ile Pro Ile Ser Gln Leu Ser Val Glu Asn Leu  
   130                          135                          140  
 Ser Asp Ala Ile Arg Phe Met Leu Gln Pro Glu Val Lys Ser Gln Ala  
   145                          150                          155                          160  
 Met Glu Met Ala Lys Leu Ile Glu Asn Glu Asp Gly Val Ala Ala Ala  
                           165                          170                          175  
 Val Asp Ala Phe His Arg His Leu Pro Glu Glu Phe Pro Ser Ser Ser  
           180                          185                          190  
 Val Ser Ser Asp Gly Glu Glu His Pro Asn Pro Phe Leu Trp Leu Phe  
           195                          200                          205  
 Leu Gln Val Glu Lys Trp Cys Cys Leu Pro Cys Ser Lys  
   210                          215                          220

<210> 246  
 <211> 114  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 246  
 Leu Asp Asn Cys Pro His Asp Trp Leu Phe Leu Arg Cys Ser Ala Val  
   1                          5                          10                          15  
 Val His His Gly Gly Ala Gly Thr Thr Ala Ala Gly Leu Lys Ala Ala  
           20                          25                          30  
 Cys Pro Thr Thr Val Val Pro Phe Phe Gly Asp Gln Pro Phe Trp Gly  
           35                          40                          45  
 Glu Arg Val His Ala Arg Gly Val Gly Pro Val Pro Ile Pro Val Asp

50                      55                      60  
 Glu Phe Ser Leu Glu Lys Leu Val Asp Ala Ile Arg Phe Met Leu Asp  
 65                      70                      75                      80  
 Pro Lys Val Lys Gln Cys Ala Glu Glu Leu Ala Lys Asp Met Glu His  
                     85                      90                      95  
 Glu Asp Gly Val Glu Gly Ala Val Lys Ala Phe Tyr Lys His Phe Pro  
                     100                      105                      110  
 Arg Glu

<210> 247  
 <211> 140  
 <212> PRT  
 <213> Pinus radiata

<400> 247  
 Met Ala Thr Gly Gly Gly Ala Leu Asp Leu Ala Ser Gly Met Gly Gly  
 1                      5                      10                      15  
 Asn Ile Glu Lys Glu Gln Met Leu Thr Ala Val Glu Glu Tyr Glu Lys  
                     20                      25                      30  
 Tyr His Met Tyr Tyr Gly Gly Asp Glu Gly Ser Arg Lys Ser Asn Tyr  
                     35                      40                      45  
 Thr Asp Met Val Asn Lys Tyr Tyr Asp Leu Ala Thr Ser Phe Tyr Glu  
 50                      55                      60  
 Tyr Gly Trp Gly Glu Ser Phe His Phe Ala His Arg Trp Lys Gly Glu  
 65                      70                      75                      80  
 Thr Leu Arg Glu Ser Ile Lys Arg His Glu His Phe Leu Ala Leu His  
                     85                      90                      95  
 Leu Cys Leu Lys Pro Ala Met Lys Val Leu Asp Val Gly Cys Gly Ile  
                     100                      105                      110  
 Gly Gly Pro Leu Arg Glu Ile Ala Arg Phe Ser Arg Thr Ser Ile Thr  
                     115                      120                      125  
 Gly Leu Asn Asn Asn Ala Tyr Gln Ile Ser Arg Gly  
                     130                      135                      140

<210> 248  
 <211> 152  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 248  
 Met Ser Lys Ala Gly Ala Met Asp Leu Ala Thr Gly Leu Gly Gly Lys  
 1                      5                      10                      15  
 Met Asp Lys Ser Asp Val Leu Ser Ala Val Asp Lys Tyr Glu Lys Tyr  
                     20                      25                      30  
 His Val Cys Tyr Gly Gly Asp Glu Glu Glu Arg Arg Ala Asn Tyr Ser  
                     35                      40                      45  
 Asp Met Val Asn Lys Tyr Tyr Asp Leu Ala Thr Ser Phe Tyr Glu Phe  
 50                      55                      60  
 Gly Trp Gly Glu Ser Phe His Phe Ala His Arg Trp Lys Gly Glu Ser  
 65                      70                      75                      80  
 Leu Arg Glu Ser Ile Lys Arg His Glu His Phe Leu Ala Leu Gln Leu  
                     85                      90                      95  
 Gly Leu Lys Pro Gly His Lys Val Leu Asp Val Gly Cys Gly Ile Gly  
                     100                      105                      110  
 Gly Pro Leu Arg Glu Ile Ala Arg Phe Ser Ser Ala Ser Val Thr Gly  
                     115                      120                      125  
 Leu Asn Asn Asn Glu Tyr Gln Ile Thr Arg Gly Lys Glu Leu Asn Arg  
                     130                      135                      140  
 Ile Ala Gly Val Asp Lys Thr Cys  
 145                      150

<210> 249  
 <211> 100  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 249  
 Met Ser Lys Ala Gly Ala Met Asp Leu Ala Thr Gly Leu Gly Gly Lys  
 1 5 10 15  
 Met Asp Lys Ser Asp Val Leu Ser Ala Val Asp Lys Tyr Glu Lys Tyr  
 20 25 30  
 His Val Cys Tyr Gly Gly Asp Glu Glu Glu Arg Arg Ala Asn Tyr Ser  
 35 40 45  
 Asp Met Val Asn Lys Tyr Tyr Asp Leu Ala Thr Ser Phe Tyr Glu Phe  
 50 55 60  
 Gly Trp Gly Glu Ser Phe His Phe Ala His Arg Trp Lys Gly Glu Ser  
 65 70 75 80  
 Leu Arg Glu Ser Ile Lys Arg His Glu His Phe Leu Ala Leu Gln Leu  
 85 90 95  
 Gly Leu Lys Pro  
 100

<210> 250  
 <211> 148  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 250  
 Ala Met Pro Trp Tyr Cys Ala Leu Pro Thr Leu Ser Glu Tyr Met Val  
 1 5 10 15  
 Glu Asn Gly Trp Thr Lys Cys Phe Ser Arg Ile Ser Asp Val Gly Trp  
 20 25 30  
 Leu Ala Tyr Leu Val Tyr Leu Ser Ile Tyr Leu Val Met Ala Glu Phe  
 35 40 45  
 Gly Ile Tyr Trp Met His Arg Glu Leu His Asp Ile Lys Pro Leu Tyr  
 50 55 60  
 Lys His Leu His Ala Thr His His Ile Tyr Asn Lys Gln Asn Thr Leu  
 65 70 75 80  
 Ser Pro Phe Ala Gly Leu Ala Phe His Pro Leu Asp Gly Ile Leu Gln  
 85 90 95  
 Ala Val Pro His Val Met Ala Leu Phe Leu Val Pro Thr His Phe Thr  
 100 105 110  
 Thr His Ile Ala Leu Leu Phe Leu Glu Ala Ile Trp Thr Ala Asn Ile  
 115 120 125  
 His Asp Cys Ile His Gly Lys Leu Trp Pro Val Met Gly Ala Gly Tyr  
 130 135 140  
 His Thr Ile His  
 145

<210> 251  
 <211> 201  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 251  
 Phe Met Ser Cys Leu Pro Asn Met Ile Val Met Ala Pro Ser Asp Glu  
 1 5 10 15  
 Asp Glu Leu Val Asp Met Val Glu Thr Ala Ala Ile Val Asp Asp Arg  
 20 25 30  
 Pro Ile Cys Phe Arg Tyr Pro Arg Gly Ala Ile Val Arg Thr Asp Lys  
 35 40 45

Ser Leu Ser Gln Gly Ile Pro Ile Glu Ile Gly Lys Gly Arg Ile Leu  
 50 55 60  
 Ala Glu Gly Lys Asp Val Ala Leu Leu Gly Tyr Gly Ser Met Val Gln  
 65 70 75 80  
 Asn Cys Val Lys Ala Arg Ser Leu Leu Ser Lys Leu Gly Ile Glu Val  
 85 90 95  
 Thr Val Ala Asp Ala Arg Phe Cys Lys Pro Leu Asp Ile Gly Leu Leu  
 100 105 110  
 Arg Glu Leu Cys Glu Asn His Ala Phe Leu Val Thr Val Glu Glu Gly  
 115 120 125  
 Ser Ile Gly Gly Phe Gly Ser His Val Ala Gln Phe Ile Ala Leu Asp  
 130 135 140  
 Gly Arg Leu Asp Gly Arg Ile Lys Trp Arg Pro Ile Val Leu Pro Asp  
 145 150 155 160  
 Ala Tyr Val Glu His Ala Ser Pro Asn Glu Gln Leu Ser Leu Ala Gly  
 165 170 175  
 Leu Thr Gly His His Ile Ala Ala Thr Val Leu Ser Leu Leu Gly Arg  
 180 185 190  
 Thr Arg Glu Ala Leu Leu Leu Met Cys  
 195 200

<210> 252  
 <211> 138  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 252  
 Asp Ile Lys Lys Ile Val Glu Leu Met Ser Asp Leu His Phe Ile Tyr  
 1 5 10 15  
 Asn Thr His Arg Phe Ala Tyr Leu Tyr Ser Lys Phe Asn Ser Ser Ile  
 20 25 30  
 Tyr Met Tyr Lys Phe Ser Leu Asp Thr Asp Leu Asn Ile Val Lys Lys  
 35 40 45  
 Met Ser Gly Phe Asp Val Glu Gly Val Cys His Ala Asp Glu Leu Phe  
 50 55 60  
 Tyr Phe Phe Ser Thr Asn Met Thr Lys Asp Tyr Tyr Glu Ser Glu Asp  
 65 70 75 80  
 Lys Ile Lys Glu Tyr Val Trp Lys Val Thr Lys Leu Trp Thr Asn Phe  
 85 90 95  
 Ala Lys Thr Ser Asn Pro Thr Pro Asp Thr Ser Leu Gly Val Ser Trp  
 100 105 110  
 Pro Arg Tyr Thr Met Ala Asn Lys Glu Tyr Leu Asp Ile Asn Thr Gln  
 115 120 125  
 Leu Thr Thr Gly Arg Tyr Ser Glu Arg Glu  
 130 135

<210> 253  
 <211> 610  
 <212> PRT  
 <213> Pinus radiata

<400> 253  
 Cys Leu Leu Leu Leu Gln Leu Lys Leu Phe Cys Ser Pro Ile Asn Met  
 1 5 10 15  
 Ala Ile Ala Ser Arg Ala Gly Val Ala Pro Ile Leu Gln Val Asp Cys  
 20 25 30  
 His Phe Thr His Phe Asn Ser Met Thr Glu Leu Gly Ser Arg Asn Ser  
 35 40 45  
 Met Met Phe Gln Ser Ala Ile Pro Cys Ser Phe Arg Gln Ile Arg Ala  
 50 55 60  
 Thr Thr Lys Arg Lys Arg Cys Val Leu Leu Ala Lys Leu Ser Asn Ser

65	70	75	80
Asp Gly Glu Asn Gly Lys Asn Val Lys Ala Ala Val Glu Ile Ala Ser			
	85	90	95
Lys Ser Gly Phe Pro Ala Glu Lys Pro Pro Thr Pro Leu Leu Asp Thr			
	100	105	110
Val Asn Tyr Pro Val His Leu Lys Asn Leu Ser Ile Gln Asp Leu Glu			
	115	120	125
Gln Leu Ala Thr Glu Ile Arg Ala Glu Leu Val Phe Gly Val Ala Lys			
	130	135	140
Thr Gly Gly His Leu Gly Gly Ser Leu Gly Val Val Asp Leu Thr Val			
	145	150	155
Ala Leu His His Val Phe Asp Ser Pro Glu Asp Arg Ile Ile Trp Asp			
	165	170	175
Val Gly His Gln Ser Tyr Pro His Lys Ile Leu Thr Gly Arg Arg Ser			
	180	185	190
Lys Met His Thr Ile Arg Gln Thr Ser Gly Leu Ala Gly Phe Pro Lys			
	195	200	205
Arg Asp Glu Ser Lys Tyr Asp Ala Phe Gly Ala Gly His Ser Ser Thr			
	210	215	220
Ser Ile Ser Ala Gly Leu Gly Met Ala Val Gly Arg Asp Leu Leu Lys			
	225	230	235
Lys Asn Asn His Val Val Ala Val Ile Gly Asp Gly Ala Met Thr Ala			
	245	250	255
Gly Gln Ala Tyr Glu Ala Met Asn Asn Ser Gly Tyr Leu Glu Ser Asn			
	260	265	270
Leu Ile Ile Ile Leu Asn Asp Asn Lys Gln Val Ser Leu Pro Thr Ala			
	275	280	285
Thr Leu Asp Gly Ala Ala Pro Val Gly Ala Leu Thr Arg Ala Leu			
	290	295	300
Thr Lys Leu Gln Ser Ser Lys Lys Leu Arg Lys Leu Arg Glu Ala Ala			
	305	310	315
Lys Gly Leu Thr Lys Gln Ile Gly Gly Pro Thr His Glu Val Ala Ser			
	325	330	335
Lys Val Asp Lys Tyr Ala Arg Gly Leu Ile Ser Pro Ala Ser Ser Ser			
	340	345	350
Leu Phe Asp Glu Leu Gly Leu Tyr Tyr Ile Gly Pro Val Asp Gly His			
	355	360	365
Asn Ile Glu Asp Met Val Thr Ile Leu Glu Lys Ile Lys Ser Met Pro			
	370	375	380
Ala Thr Gly Pro Val Leu Ile His Leu Val Thr Glu Lys Gly Lys Gly			
	385	390	395
Tyr Pro Pro Ala Glu Glu Ala Ala Asp Lys Leu His Gly Val Val Lys			
	405	410	415
Phe Asp Pro Val Thr Gly Lys Gln Phe Lys Ser Lys Ser Ser Val Leu			
	420	425	430
Ser Tyr Thr Gln Tyr Phe Ala Glu Ala Leu Ile Ala Glu Ala Glu Val			
	435	440	445
Asp Ser Lys Ile Val Ala Ile His Ala Ala Met Gly Gly Gly Thr Gly			
	450	455	460
Leu Asn Tyr Phe Gln Lys Lys Phe Pro Glu Arg Cys Phe Asp Val Gly			
	465	470	475
Ile Ala Glu Gln His Ala Val Thr Phe Ala Ala Gly Leu Ala Thr Glu			
	485	490	495
Gly Leu Lys Pro Phe Cys Ala Ile Tyr Ser Thr Phe Leu Gln Arg Gly			
	500	505	510
Tyr Asp Gln Val Val His Asp Val Asp Leu Gln Lys Leu Pro Val Arg			
	515	520	525
Phe Ala Met Asp Arg Ala Gly Leu Val Gly Ala Asp Gly Pro Thr His			
	530	535	540
Cys Gly Ser Phe Asp Val Ala Tyr Met Ala Cys Leu Pro Asn Met Ile			
545	550	555	560

```
<210> 254
<211> 147
<212> PRT
<213> Eucalyptus grandis
```

```
<210> 255
<211> 123
<212> PRT
<213> Eucalyptus grandis
```

```
<210> 256
<211> 127
```

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 256

```

Arg Pro Cys His Leu Glu Trp Ile His Ile His Lys Thr Ala Val Ile
 1           5           10           15
Leu Glu Cys Ser Val Val Cys Gly Asp Ile Ile Ser Gly Ala Ser Glu
          20           25           30
Asn Glu Ile Glu Arg Ile Lys Ser Tyr Ala Arg Ser Val Gly Leu Leu
          35           40           45
Phe Gln Val Val Asp Asp Ile Leu Asp Val Thr Lys Ser Ser Lys Glu
          50           55           60
Leu Gly Lys Thr Ala Gly Lys Asp Leu Ile Thr Asp Lys Ala Thr Tyr
65           70           75           80
Pro Lys Leu Met Gly Leu Glu Thr Ala Lys Gln Phe Ala Val Glu Leu
          85           90           95
Leu Gly Arg Ala Lys Glu Asp Leu Ser Cys Phe Asp Pro Lys Lys Ala
          100          105          110
Ala Pro Leu Leu Gly Ile Ala Glu Tyr Ile Ala Phe Arg Gln Asn
          115          120          125

```

&lt;210&gt; 257

&lt;211&gt; 196

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 257

```

Ala Cys Ala Val Glu Met Ile His Thr Met Ser Leu Ile His Asp Asp
 1           5           10           15
Leu Pro Cys Met Asp Asn Asp Asp Leu Arg Arg Gly Lys Pro Thr Asn
          20           25           30
His Lys Val Tyr Gly Glu Asp Val Ala Val Leu Ala Gly Asp Ala Leu
          35           40           45
Leu Ala Tyr Ala Phe Glu His Ile Ala Val Glu Thr Lys Gly Val Ser
          50           55           60
Pro Thr Arg Ile Val Arg Ala Ile Phe Glu Leu Ala Arg Ser Ile Gly
65           70           75           80
Ala Glu Gly Leu Val Ala Gly Gln Val Val Asp Ile Ser Ser Glu Gly
          85           90           95
Ile Ala Asn Val Gly Leu Glu His Leu Glu Phe Ile His Leu His Lys
          100          105          110
Thr Ala Ala Leu Leu Glu Ala Ser Val Val Leu Gly Ala Ile Met Gly
          115          120          125
Gly Gly Ser Asn Glu Glu Val Lys Leu Arg Gly Phe Ala Arg Cys
          130          135          140
Ile Gly Leu Leu Phe Gln Val Val Asp Asp Ile Leu Asp Leu Thr Gln
145          150          155          160
Ser Ser Gln Glu Leu Gly Lys Thr Ala Gly Lys Asp Leu Val Ala Asp
          165          170          175
Lys Val Thr Tyr Pro Lys Leu Met Gly Ile Glu Lys Ser Arg Glu Leu
          180          185          190
Ala Asn Lys Leu
          195

```

&lt;210&gt; 258

&lt;211&gt; 159

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 258

```

Met Gly Ser Leu Gly Ala Ile Leu Lys His Pro Asp Asp Phe Tyr Pro

```



1	5	10	15
Leu Leu Lys	Leu Lys Ile Ala Ala	Arg Asn Ala Glu Lys Arg Ile Pro	
	20	25	30
Pro Gln Pro	His Trp Gly Phe Cys Tyr Ser Met Leu His Lys Val Ser		
	35	40	45
Arg Ser Phe	Gly Leu Val Ile Gln Gln Leu Gly Pro Glu Leu Arg Asp		
	50	55	60
Ala Val Cys	Ile Phe Tyr Leu Val Leu Arg Ala Leu Asp Thr Val Glu		
65	70	75	80
Asp Asp Thr	Ser Ile Pro Thr Asp Val Lys Val Pro Ile Leu Lys Ala		
	85	90	95
Phe His Gln	His Val Tyr Asp Lys Glu Trp His Phe Ser Cys Gly Thr		
	100	105	110
Lys Glu Tyr	Lys Val Leu Met Asp Gln Phe His His Val Ser Thr Ala		
	115	120	125
Phe Leu Glu	Leu Gly Lys Ser Tyr Gln Glu Ala Ile Asp Asp Ile Thr		
	130	135	140
Lys Arg Met	Gly Ala Gly Met Ala Lys Phe Ile Cys Gln Glu Val		
145	150	155	

&lt;210&gt; 259

&lt;211&gt; 106

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 259

Met Ala Ile	Tyr Thr Pro Gln Pro Ala His Arg Leu Ile Ser Trp Ser
1	5 10 15
Thr Met Glu	Asn His Thr Val Ala Ile Ala Val Ala Ile Gly Phe Val
	20 25 30
Ser Val Leu	Leu Ser Tyr Tyr Ile Val Leu Asn Arg Trp Lys Arg Arg
	35 40 45
Ser Asn Gly	Leu Arg Gly Ile Gln Ser Lys Ser Phe Glu Lys Ser Thr
	50 55 60
Asp Asp Asn	Gly Ile Ala Ile Glu Ala Ala Gly Gly Thr Asp Val Ile
65	70 75 80
Ile Val Gly	Ala Gly Val Ala Gly Ser Ala Leu Ala Tyr Thr Leu Gly
	85 90 95
Lys Asp Gly	Arg Arg Ile His Val Ile Glu
	100 105

&lt;210&gt; 260

&lt;211&gt; 93

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 260

Met Ala Ile	Tyr Thr Pro Gln Pro Ala His Arg Leu Ile Ser Trp Ser
1	5 10 15
Thr Met Glu	Asn His Thr Val Val Ile Ala Ala Ala Ile Ser Phe Val
	20 25 30
Ser Val Leu	Leu Ser Tyr Tyr Ile Val Leu Ser Arg Trp Lys Arg Arg
	35 40 45
Ser Asn Gly	Leu Arg Gly Ile Gln Ser Lys Ser Phe Glu Lys Ser Thr
	50 55 60
Asp Asp Asn	Gly Ile Ala Ile Glu Ala Ala Gly Gly Thr Asp Val Ile
65	70 75 80
Ile Val Gly	Ala Gly Val Ala Gly Ser Ala Leu Ala Tyr
	85 90

&lt;210&gt; 261

<211> 217  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 261  
 Pro Gln Leu Tyr Lys Ala Phe Ile Ala Ala Ile Asp Lys Gly Asn Ile  
 1 5 10 15  
 Lys Ser Met Pro Asn Arg Ser Met Pro Ala Asn Pro Gln Pro Thr Pro  
 20 25 30  
 Gly Ala Leu Leu Met Gly Asp Ala Phe Asn Met Arg His Pro Leu Thr  
 35 40 45  
 Gly Gly Gly Met Thr Val Ala Leu Ser Asp Ile Val Leu Leu Arg Asn  
 50 55 60  
 Leu Leu Arg Pro Leu Gln Asp Leu Asn Asp Ala Ser Ala Leu Cys Lys  
 65 70 75 80  
 Tyr Leu Glu Ser Phe Tyr Thr Leu Arg Lys Pro Val Ala Ser Thr Ile  
 85 90 95  
 Asn Thr Leu Ala Gly Ala Leu Tyr Lys Val Phe Cys Ala Ser Pro Asp  
 100 105 110  
 Pro Ala Arg Lys Glu Met Arg Gln Ala Cys Phe Asp Tyr Leu Ser Leu  
 115 120 125  
 Gly Gly Leu Cys Ser Thr Gly Pro Val Ser Leu Leu Ser Gly Leu Asn  
 130 135 140  
 Pro Arg Pro Met His Leu Val Cys His Phe Phe Ala Val Ala Val Tyr  
 145 150 155 160  
 Gly Val Gly Arg Leu Cys Leu Pro Phe Pro Ser Pro Lys Arg Met Trp  
 165 170 175  
 Leu Gly Ala Arg Leu Val Lys Gly Ala Ser Gly Ile Ile Phe Pro Ile  
 180 185 190  
 Ile Arg Asp Glu Gly Val Arg Gln Met Phe Phe Pro Ala Thr Val Pro  
 195 200 205  
 Ala Tyr His Arg Ala Pro Pro Val His  
 210 215

<210> 262  
 <211> 94  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 262  
 Met Glu Asp Asp Arg Asp Arg Gly Leu Leu Tyr Asp Ser Asp Pro Ser  
 1 5 10 15  
 Ser Ser Ser Leu Ser Pro Pro Arg Pro Phe Ala Leu Thr Phe Phe Asp  
 20 25 30  
 Arg Glu Arg His Val Thr Phe Leu Glu Met Met Tyr His Met Leu Pro  
 35 40 45  
 Arg Pro Tyr Gln Ser Gln Glu Ile Asn His Leu Thr Leu Ala Tyr Phe  
 50 55 60  
 Val Ile Ser Gly Leu Asp Ile Leu Asp Ala Leu Asp Arg Val His Lys  
 65 70 75 80  
 Asp Ala Val Ala Asp Trp Val Leu Ser Phe Gln Ala His Phe  
 85 90

<210> 263  
 <211> 81  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 263  
 Glu Ile Leu Thr Lys Val Ile Ser Leu Ala Ser Ile Met Asp Asp Ile  
 1 5 10 15

Tyr Asp Val Tyr Gly Thr Leu Glu Glu Leu Ala Leu Leu Asn Glu Ala  
                   20                  25                  30  
 Ile Gln Lys Trp Asp Phe Asp Ala Met Asp Gly Leu Pro Glu Tyr Met  
                   35                  40                  45  
 Gln Ala Tyr Phe Lys Glu Phe Leu Gln Leu Tyr Glu Tyr Ile Gly Asn  
                   50                  55                  60  
 Gln Leu Ala Ala Lys Gly Arg Ser Tyr Arg Leu Ile Tyr Ala Lys Glu  
                   65                  70                  75                  80  
 Val

<210> 264  
 <211> 125  
 <212> PRT  
 <213> Pinus radiata

<400> 264  
 Leu Tyr Arg Ala Ser Leu Ile Ala Phe Pro Gly Glu Lys Val Met Asp  
   1                  5                  10                  15  
 Glu Ala Glu Thr Phe Ser Ala Lys Tyr Leu Lys Glu Ala Leu Gln Lys  
                   20                  25                  30  
 Ile Pro Val Ser Ser Leu Ser Arg Glu Ile Gly Asp Val Leu Glu Tyr  
                   35                  40                  45  
 Gly Trp His Thr Tyr Leu Pro Arg Leu Glu Ala Arg Asn Tyr Ile Asp  
                   50                  55                  60  
 Val Phe Gly Gln Asp Thr Glu Asn Ser Lys Ser Tyr Met Lys Thr Glu  
                   65                  70                  75                  80  
 Lys Leu Leu Glu Leu Ala Lys Leu Glu Phe Asn Ile Phe His Ala Leu  
                                   85                  90                  95  
 Gln Lys Arg Glu Leu Glu Tyr Leu Val Arg Trp Trp Lys Gly Ser Gly  
                   100                  105                  110  
 Ser Pro Gln Met Thr Phe Cys Arg His Arg His Val Glu  
                   115                  120                  125

<210> 265  
 <211> 219  
 <212> PRT  
 <213> Pinus radiata

<400> 265  
 Met Pro Gln Asp Met Lys Ile Cys Phe Lys Gly Phe Tyr Asn Thr Phe  
   1                  5                  10                  15  
 Asn Glu Ile Ala Glu Glu Gly Arg Lys Arg Gln Gly Arg Asp Val Leu  
                   20                  25                  30  
 Ser Tyr Ile Gln Lys Val Trp Glu Val Gln Leu Glu Ala Tyr Thr Lys  
                   35                  40                  45  
 Glu Ala Glu Trp Ser Ala Val Arg Tyr Val Pro Ser Tyr Asp Glu Tyr  
                   50                  55                  60  
 Ile Gly Asn Ala Ser Val Ser Ile Ala Leu Gly Thr Val Val Leu Ile  
                   65                  70                  75                  80  
 Ser Ala Leu Phe Thr Gly Glu Ile Leu Thr Asp Asp Ile Leu Ser Lys  
                   85                  90                  95  
 Ile Gly Arg Asp Ser Arg Phe Leu Tyr Leu Met Gly Leu Thr Gly Arg  
                   100                  105                  110  
 Leu Val Asn Asp Thr Lys Thr Tyr Gln Ala Glu Arg Gly Gln Gly Glu  
                   115                  120                  125  
 Val Ala Ser Ala Val Gln Cys Tyr Met Lys Asp His Pro Glu Ile Ser  
                   130                  135                  140  
 Glu Glu Glu Ala Leu Lys His Val Tyr Thr Ile Met Asp Asn Ala Leu  
                   145                  150                  155                  160  
 Asp Glu Leu Asn Arg Glu Phe Val Asn Asn Arg Asp Val Pro Asp Thr

165 170 175  
 Cys Arg Arg Leu Val Phe Glu Thr Ala Arg Ile Met Gln Leu Phe Tyr  
 180 185 190  
 Met Asp Gly Asp Gly Leu Thr Leu Ser His Asn Met Glu Ile Lys Glu  
 195 200 205  
 His Val Lys Asn Cys Leu Phe Gln Pro Val Ala  
 210 215

<210> 266  
 <211> 423  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 266  
 Leu Asp Cys Glu Pro Val Val Gln Lys Pro Lys Leu Val Asp Pro Val  
 1 5 10 15  
 Val Gln Asp Ala Pro Lys Glu Lys Val Ile Glu Ala Val Pro Ser Ala  
 20 25 30  
 Met Pro Glu Glu Asp Glu Glu Ile Ile Lys Ser Val Val Glu Gly Lys  
 35 40 45  
 Met Pro Ser Tyr Ser Leu Glu Ser Lys Leu Gly Asp Cys Lys Arg Ala  
 50 55 60  
 Ala Ala Ile Arg Arg Glu Ala Leu Gln Arg Ile Thr Gly Lys Ser Leu  
 65 70 75 80  
 Ser Gly Leu Pro Leu Glu Gly Phe Asp Tyr Glu Ser Ile Leu Gly Gln  
 85 90 95  
 Cys Cys Glu Met Pro Val Gly Tyr Val Gln Ile Pro Val Gly Ile Ala  
 100 105 110  
 Gly Pro Leu Leu Leu Asp Gly Arg Glu Tyr Ser Val Pro Met Ala Thr  
 115 120 125  
 Thr Glu Gly Cys Leu Val Ala Ser Thr Asn Arg Gly Cys Lys Ala Ile  
 130 135 140  
 Phe Val Ser Gly Gly Ala Thr Ser Val Leu Leu Arg Asp Gly Met Thr  
 145 150 155 160  
 Arg Ala Pro Ile Val Arg Phe Gly Thr Ala Lys Arg Ala Ala Asp Leu  
 165 170 175  
 Lys Phe Phe Val Glu Asn Pro Ala Asn Phe Glu Ser Leu Ala Val Ile  
 180 185 190  
 Phe Asn Arg Ser Ser Arg Phe Ala Arg Leu Gln Ser Ile Lys Cys Ala  
 195 200 205  
 Ile Ala Gly Lys Asn Leu Tyr Met Arg Phe Ser Cys Ser Thr Gly Asp  
 210 215 220  
 Ala Met Gly Met Asn Met Val Ser Lys Gly Val Gln Asn Val Leu Asp  
 225 230 235 240  
 Phe Leu Gln Ser Asp Phe Pro Asp Met Asp Val Leu Gly Ile Ser Gly  
 245 250 255  
 Asn Phe Cys Ala Asp Lys Lys Pro Ala Ala Val Asn Trp Ile Glu Gly  
 260 265 270  
 Arg Gly Lys Ser Val Val Cys Glu Ala Thr Ile Lys Gly Asp Val Val  
 275 280 285  
 Arg Lys Val Leu Lys Thr Ser Val Glu Ala Leu Val Glu Leu Asn Met  
 290 295 300  
 Leu Lys Asn Leu Thr Gly Ser Ala Met Ala Gly Ala Leu Gly Gly Phe  
 305 310 315 320  
 Asn Ala His Ala Ser Asn Ile Val Ala Ala Ile Phe Ile Ala Thr Gly  
 325 330 335  
 Gln Asp Pro Ala Gln Asn Val Glu Ser Ser His Cys Ile Thr Met Met  
 340 345 350  
 Glu Ala Ile Asn Asp Gly Lys Asp Leu His Val Ser Val Thr Met Pro  
 355 360 365  
 Ser Ile Glu Val Gly Thr Val Gly Gly Gly Thr Gln Leu Ala Ser Gln

370                      375                      380  
 Ser Ala Cys Leu Asn Leu Leu Gly Val Lys Gly Ala Asn Lys Glu Leu  
 385                      390                      395                      400  
 Ala Gly Ala Asn Ser Arg Leu Leu Ala Thr Val Val Ser Gly Ala Val  
                     405                      410                      415  
 Leu Ala Ala Glu Leu Ser Ser  
                     420

<210> 267  
 <211> 112  
 <212> PRT  
 <213> Pinus radiata

<400> 267  
 Met Ser Leu Ile Ser Ala Val Pro Leu Ala Ser Ser Cys Val Ser Lys  
 1                      5                      10                      15  
 Ser Leu Ile Ser Ser Val Arg Glu His Lys Ala Leu Arg Arg Ala Ile  
                     20                      25                      30  
 Ala Thr Leu Gln Met Ser Arg Pro Gly Lys Ser Val Ala Ala Ser Thr  
                     35                      40                      45  
 Arg Met Ser Ser Ala Thr Ala Gly Ser Asp Asp Gly Val Lys Arg Arg  
                     50                      55                      60  
 Ile Gly Asp Tyr His Ser Asn Leu Trp Glu Asp Asn Phe Ile Gln Ser  
 65                      70                      75                      80  
 Leu Ser Ser Pro Tyr Gly Ala Ser Ser Tyr Gly Glu His Ala Asp Arg  
                     85                      90                      95  
 Leu Ile Gly Glu Val Lys Gly Ile Phe Asn Ser Phe Ser Ile Ala Asp  
                     100                      105                      110

<210> 268  
 <211> 165  
 <212> PRT  
 <213> Pinus radiata

<400> 268  
 Met Ser Leu Ile Ser Ala Val Pro Leu Ala Ser Ser Ser Val Ser Lys  
 1                      5                      10                      15  
 Ser Leu Ile Ser Ser Val Arg Glu His Lys Ala Leu Arg Arg Ala Ile  
                     20                      25                      30  
 Ala Thr Leu Gln Met Ser Arg Pro Gly Lys Ser Val Ala Ala Ser Thr  
                     35                      40                      45  
 Lys Met Ser Ser Ala Thr Ala Gly Ser Asp Asp Gly Val Lys Arg Arg  
                     50                      55                      60  
 Ile Gly Asp Tyr His Ser Asn Leu Trp Asp Asp Asn Val Ile Gln Ser  
 65                      70                      75                      80  
 Leu Ser Ser Pro Tyr Gly Ala Ser Ser Tyr Gly Glu His Ala Asp Arg  
                     85                      90                      95  
 Leu Ile Gly Glu Val Lys Glu Ile Phe Asn Ser Phe Ser Ile Ala Asp  
                     100                      105                      110  
 Gly Glu Leu Thr Ser Pro Val Asn Asp Leu Leu Gln Gln Leu Trp Met  
                     115                      120                      125  
 Val Asp Asn Val Glu Arg Leu Gly Ile Asp Arg His Phe Gln Thr Glu  
                     130                      135                      140  
 Ile Lys Val Ala Leu Asp Tyr Gly Tyr Arg Tyr Trp Ser Glu Lys Gly  
 145                      150                      155                      160  
 Ile Glu Cys Gly Glu  
                     165

<210> 269  
 <211> 144  
 <212> PRT

## &lt;213&gt; Pinus radiata

&lt;400&gt; 269

```

Ser Thr Leu Gln Leu Ser Arg Arg Gly Lys Pro Val Thr Ala Cys Lys
 1          5          10          15
Lys Val Ser Leu Ser Thr Ala Val Ser Asp Asp Gly Ala Lys Arg Arg
          20          25          30
Val Gly Asp His His Ser Asn Leu Trp Asp Asp Asn Phe Ile Lys Ser
          35          40          45
Leu Ser Ser Pro Tyr Gly Ala Ser Ser Tyr Arg Glu His Ala Asp Arg
          50          55          60
Val Ile Gly Glu Val Lys Glu Ile Phe Asn Ser Leu Ser Met Thr Asp
65          70          75          80
Gly Glu Leu Ile Ser Pro Asp Asn Asp Leu Leu Gln Arg Leu Ser Met
          85          90          95
Val Asp Asn Ile Glu Arg Leu Gly Ile Asp Arg His Phe Gln Thr Glu
          100          105          110
Ile Lys Leu Thr Leu Asp Tyr Val Tyr Ser Tyr Trp Ser Glu Lys Gly
          115          120          125
Ile Gly Tyr Gly Arg Glu Ser Ala Ile Thr Asp Leu Asn Thr Thr Ser
130          135          140

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&lt;210&gt; 270

&lt;211&gt; 106

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 270

```

Gly Thr Lys Ala Lys Gly Asn Lys Gln Leu Gln Asn Asn Val Ile Lys
 1          5          10          15
Val Ile Cys Asn Thr Asp Lys Ser Arg Gly Phe Asn Val Leu Arg Asp
          20          25          30
Val Ser Met Pro Gln Ile Met Ile Lys Ser Cys Lys Val Ser Pro Asp
          35          40          45
Ala Arg Pro Tyr Gln Asn Leu Gly Gly Pro Ala Ser Ser Glu Arg Pro
          50          55          60
Phe Leu Ala Phe Phe Ala Gly Gln Met His Gly Thr Leu Arg Pro Glu
65          70          75          80
Ile Leu Lys His Trp Gly Asn Glu Thr Asp Pro Asn Met Lys Ile Phe
          85          90          95
Ala Val Gly Gln Ser His Pro Gly Ser Leu
          100          105

```

&lt;210&gt; 271

&lt;211&gt; 169

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 271

```

Lys Ala Arg Ala Val Trp Glu Asn Phe Lys Asp Asn Pro Leu Phe Asp
 1          5          10          15
Ile Ser Thr Asp His Pro Thr Thr Tyr Tyr Glu Asp Met Gln Arg Ala
          20          25          30
Val Phe Cys Leu Cys Pro Leu Gly Trp Ala Pro Trp Ser Pro Arg Leu
          35          40          45
Val Glu Ala Val Val Phe Gly Cys Ile Pro Val Ile Ile Ala Asp Asp
          50          55          60
Ile Val Leu Pro Phe Ala Asp Ala Ile Pro Trp Glu Glu Ile Gly Val
65          70          75          80
Phe Val Ala Glu Glu Asp Val Pro Ser Leu Asp Thr Ile Leu Thr Ser
          85          90          95

```

Ile Ser Pro Glu Val Ile Leu Arg Lys Gln Arg Leu Leu Ala Asn Pro  
 100 105 110  
 Ser Met Lys Arg Ala Met Leu Phe Pro Gln Pro Ala Gln Ser Gly Asp  
 115 120 125  
 Ala Phe His Gln Ile Leu Asn Gly Leu Ala Arg Lys Leu Pro His His  
 130 135 140  
 Arg Ser Val Tyr Leu Lys Pro Gly Glu Lys Val Leu Asn Trp Thr Ala  
 145 150 155 160  
 Gly Pro Val Gly Asp Leu Lys Pro Trp  
 165

&lt;210&gt; 272

&lt;211&gt; 146

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 272

Met Ser Gln Val Ser Ala Thr Pro Cys Ala Pro Pro Asn Lys Glu Thr  
 1 5 10 15  
 Gly His Val Ile Glu Arg Arg Ser Ala Gly Tyr His Pro Ser Val Trp  
 20 25 30  
 Gly Asp Tyr Phe Leu Lys Tyr Asp Ser Pro Ser Asn Ser Val Lys Phe  
 35 40 45  
 Lys Phe Leu Gly Arg Val Glu Gly Gln Ile Glu Glu Leu Lys Gly Glu  
 50 55 60  
 Val Lys Lys Met Leu Ile Asp Val Val Asp Lys Pro Leu Pro Lys Leu  
 65 70 75 80  
 His Leu Ile Asp Gln Ile Gln Arg Leu Gly Ile Glu Tyr His Phe Glu  
 85 90 95  
 Arg Glu Val Asp Glu Gln Leu Glu Gln Ile His Lys Ser Tyr Ser Arg  
 100 105 110  
 Leu Asp His Glu Asp Phe Lys Val Asp Asp Leu His Met Val Ala Leu  
 115 120 125  
 Ile Phe Arg Leu Leu Arg Gln His Gly Tyr Asn Ile Ser Ser Glu Ile  
 130 135 140  
 Phe Asp  
 145

&lt;210&gt; 273

&lt;211&gt; 132

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 273

Lys Lys Met Leu Ile Asp Ala Val Asp Lys Pro Leu Pro Lys Leu His  
 1 5 10 15  
 Leu Ile Asp Gln Ile Gln Arg Leu Gly Ile Glu Tyr His Phe Glu Arg  
 20 25 30  
 Glu Val Asp Glu Gln Leu Glu Gln Ile His Lys Ser Tyr Ser Arg Leu  
 35 40 45  
 Asp His Glu Asp Phe Lys Val Asp Asp Leu His Thr Val Ala Leu Ile  
 50 55 60  
 Phe Arg Leu Leu Arg Gln His Gly Tyr Asn Ile Ser Ser Glu Val Phe  
 65 70 75 80  
 Asp Lys Phe Lys Ile Ala Thr Gly Thr Ser Glu Ser Arg Leu Ile Ser  
 85 90 95  
 Asp Val Arg Gly Leu Leu Ser Leu Tyr Glu Ala Cys His Leu Arg Cys  
 100 105 110  
 His Gly Asp Ser Ile Leu Asp Glu Ala Leu Pro Phe Ala Thr Thr His  
 115 120 125  
 Leu Glu Ser Ile

130

<210> 274  
 <211> 116  
 <212> PRT  
 <213> Eucalyptus grandis

&lt;400&gt; 274

Met	Ser	Gln	Val	Ser	Ala	Thr	Pro	Cys	Ala	Pro	Ser	Asn	Lys	Gly	Thr
1				5					10					15	
Gly	His	Val	Ile	Glu	Arg	Arg	Ser	Ala	Gly	Tyr	His	Pro	Ser	Val	Trp
		20					25					30			
Gly	Asp	Tyr	Phe	Leu	Lys	Tyr	Asp	Ser	Pro	Ser	Asn	Ser	Val	Lys	Phe
	35					40					45				
Lys	Phe	Leu	Gly	Arg	Val	Glu	Gly	Gln	Ile	Glu	Glu	Leu	Lys	Gly	Glu
	50				55					60					
Val	Lys	Lys	Met	Leu	Thr	Asp	Ile	Met	Asp	Lys	Pro	Leu	Gln	Lys	Leu
65				70					75					80	
His	Leu	Ile	Asp	Gln	Ile	Gln	Arg	Leu	Gly	Ile	Glu	Tyr	His	Phe	Glu
		85							90					95	
Arg	Glu	Ile	Asp	Glu	Gln	Leu	Glu	Gln	Ile	His	Lys	Ser	Tyr	Ser	Arg
	100						105						110		
Leu	Asp	His	Glu												
		115													

<210> 275  
 <211> 214  
 <212> PRT  
 <213> Pinus radiata

&lt;400&gt; 275

Met	Ala	Thr	Phe	Ser	Asp	Glu	Thr	Pro	Val	Ser	Ser	Leu	Ala	Cys	Gly
1				5					10					15	
Leu	Ser	Ser	Asn	Ser	Gly	Leu	Ile	Arg	Arg	Thr	Ala	Asn	Pro	His	Pro
			20					25					30		
Asn	Val	Trp	Gly	Tyr	Glu	Phe	Val	Asn	Ser	Leu	Lys	Ser	Pro	Tyr	Ala
	35					40					45				
Asn	Ser	Ser	Tyr	Arg	Glu	Arg	Ala	Glu	Thr	Leu	Val	Ser	Glu	Ile	Lys
	50				55					60					
Ala	Met	Leu	Asn	Thr	Ala	Ile	Ala	Gly	Asp	Gly	Asp	Leu	Met	Ile	Thr
65				70					75					80	
Pro	Ser	Ala	Tyr	Asp	Thr	Ala	Trp	Ile	Ala	Arg	Val	Pro	Ala	Ile	Asp
			85					90					95		
Gly	Ser	Pro	Arg	Pro	Gln	Phe	Pro	Gln	Thr	Val	Asp	Trp	Ile	Leu	Lys
		100					105						110		
Asn	Gln	Leu	Lys	Asp	Gly	Ser	Trp	Gly	Thr	Gln	Ser	His	Phe	Leu	Leu
		115					120					125			
Ser	Asp	Arg	Leu	Leu	Ala	Thr	Leu	Ser	Cys	Val	Leu	Ala	Leu	Leu	Lys
	130					135					140				
Trp	Lys	Val	Gly	Asp	Ala	Gln	Val	Gln	Gln	Gly	Ile	Lys	Phe	Ile	Arg
145				150					155					160	
Ser	Asn	Leu	Leu	Lys	Asp	Glu	Ser	Asp	Glu	Asp	Ser	Leu	Val	Thr	Asp
			165					170					175		
Phe	Glu	Val	Asn	Phe	Pro	Phe	Leu	Leu	Arg	Glu	Ala	Gln	Ser	Phe	Gln
		180					185					190			
Leu	Glu	Leu	Pro	Tyr	Asp	Leu	Pro	Tyr	Ile	His	Lys	Leu	Gln	Met	Lys
		195				200						205			
Arg	Gln	Glu	Arg	Leu	Ala										
		210													

&lt;210&gt; 276



&lt;211&gt; 462

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 276

Arg Asp Ser Ala Phe Thr Asp Leu Asn Thr Thr Ala Leu Gly Phe Arg  
 1 5 10 15  
 Ile Phe Arg Leu His Gly Tyr Thr Val Ser Ser Asp Ala Phe Glu His  
 20 25 30  
 Phe Lys Asp Gln Met Gly Gln Phe Ser Ala Ser Ala Asn Asp Thr Glu  
 35 40 45  
 Leu Gln Ile Arg Ser Val Phe Asn Leu Phe Arg Ala Ser Leu Ile Ala  
 50 55 60  
 Phe Pro Glu Glu Lys Val Leu Glu Glu Ala Glu Asn Phe Ala Ala Ala  
 65 70 75 80  
 Tyr Leu Lys Ala Ala Leu Gln Thr Leu Pro Val Ser Gly Leu Ser Arg  
 85 90 95  
 Glu Ile Gln Tyr Val Phe Asp Tyr Arg Trp His Ser Asn Leu Pro Arg  
 100 105 110  
 Leu Glu Ala Arg Ser Tyr Val Asp Ile Leu Ala Asp Asn Thr Ile Ser  
 115 120 125  
 Gly Thr Pro Asp Ala Asn Thr Lys Lys Leu Leu Glu Leu Ala Lys Leu  
 130 135 140  
 Glu Phe Asn Ile Phe His Ser Leu Gln Gln Lys Glu Leu Gln Cys Leu  
 145 150 155 160  
 Trp Arg Trp Trp Lys Glu Trp Gly Cys Pro Glu Leu Thr Phe Val Arg  
 165 170 175  
 His Arg Tyr Val Glu Phe Tyr Thr Leu Val Ser Gly Thr Asp Met Val  
 180 185 190  
 Pro Glu His Ala Ala Phe Arg Leu Ser Phe Val Lys Thr Cys His Leu  
 195 200 205  
 Ile Thr Ile Leu Asp Asp Met Tyr Asp Thr Phe Gly Thr Ile Asp Glu  
 210 215 220  
 Leu Arg Leu Phe Thr Ala Ala Val Lys Arg Trp Asp Pro Ser Ala Thr  
 225 230 235 240  
 Glu Cys Leu Pro Glu Tyr Met Lys Gly Val Tyr Met Val Leu Tyr Glu  
 245 250 255  
 Thr Val Asn Glu Met Ala Lys Glu Ala Gln Lys Ser Gln Gly Arg Asp  
 260 265 270  
 Thr Leu Gly Tyr Val Arg Gln Ala Leu Glu Asp Tyr Ile Gly Ser Tyr  
 275 280 285  
 Leu Lys Glu Ala Glu Trp Ile Ala Thr Gly Tyr Val Pro Thr Phe Gln  
 290 295 300  
 Glu Tyr Phe Glu Asn Gly Lys Leu Ser Ser Gly His Arg Ile Ala Thr  
 305 310 315 320  
 Leu Gln Pro Ile Leu Thr Leu Ser Ile Pro Phe Pro His His Ile Leu  
 325 330 335  
 Gln Glu Ile Asp Phe Pro Ser Lys Phe Asn Asp Tyr Ala Cys Ser Ile  
 340 345 350  
 Leu Arg Leu Arg Gly Asp Thr Arg Cys Tyr Lys Ala Asp Ser Ala Arg  
 355 360 365  
 Gly Glu Glu Ala Ser Cys Ile Ser Cys Tyr Met Lys Glu Asn Pro Gly  
 370 375 380  
 Ser Thr Gln Glu Asp Ala Leu His His Ile Asn Gly Met Ile Glu Asp  
 385 390 395 400  
 Met Ile Lys Lys Leu Asn Trp Glu Phe Leu Lys Pro Asp Asn Asn Ala  
 405 410 415  
 Pro Ile Ser Ser Lys Lys Asn Ala Phe Asn Ile Ser Arg Gly Leu His  
 420 425 430  
 His Phe Tyr Asn Tyr Arg Asp Gly Tyr Ser Val Ala Ser Asn Glu Thr  
 435 440 445

Lys Asp Leu Val Ile Lys Thr Val Leu Glu Pro Val Leu Met  
 450 455 460

<210> 277  
 <211> 98  
 <212> PRT  
 <213> Pinus radiata

<400> 277  
 Leu Gly Glu Asp Ser Leu Thr Gly Thr Pro Asp Val Asn Thr Lys Lys  
 1 5 10 15  
 Leu Leu Glu Leu Ser Lys Leu Glu Phe Asn Ile Phe His Ser Val Gln  
 20 25 30  
 Gln Lys Glu Leu Gln Cys Leu Ser Arg Trp Trp Lys Glu Ser Gly Ser  
 35 40 45  
 Pro Glu Leu Thr Phe Ala Arg His Arg Tyr Val Glu Phe Tyr Thr Leu  
 50 55 60  
 Val Cys Gly Ile Asp Met Glu Pro Lys Asp Ala Ala Phe Arg Leu Ser  
 65 70 75 80  
 Phe Val Lys Met Cys His Leu Ile Thr Ile Leu Asp Asp Ile Tyr Asp  
 85 90 95  
 Thr Phe

<210> 278  
 <211> 63  
 <212> PRT  
 <213> Pinus radiata

<400> 278  
 Thr Glu Cys Leu Pro Glu Tyr Met Lys Gly Val Tyr Met Val Leu Tyr  
 1 5 10 15  
 Glu Thr Val Asn Glu Met Ala Lys Glu Ala Gln Lys Ser Gln Gly Arg  
 20 25 30  
 Asp Thr Leu Gly Tyr Val Arg Gln Ala Val Ile Thr Ile Asp Met Leu  
 35 40 45  
 Cys Ile Tyr Leu Asn Lys Gln Ile Leu Val Gly His Leu Phe Tyr  
 50 55 60

<210> 279  
 <211> 124  
 <212> PRT  
 <213> Pinus radiata

<400> 279  
 Ala Asp Leu Leu Asp Glu Cys Gly Pro Leu Leu Lys Lys Ala His Ala  
 1 5 10 15  
 Phe Leu Glu Lys Ser Gln Val Gln Glu Asn Ser Pro Gly Glu Phe Ser  
 20 25 30  
 Thr Trp Tyr Arg His Ile Ser Lys Gly Ala Trp Thr Leu Ser Thr Arg  
 35 40 45  
 Asp His Gly Trp Val Val Ala Asp Cys Ser Ala Glu Gly Leu Lys Ala  
 50 55 60  
 Ala Leu Glu Leu Ser Gln Leu Pro Glu Asn Ile Val Gly Lys Pro Leu  
 65 70 75 80  
 Pro Gln Gln Arg Leu Phe Ala Cys Val Asn Tyr Leu Leu Ser Met Gln  
 85 90 95  
 Asn Thr Asp Gly Gly Tyr Ala Thr Tyr Asp Leu Thr Arg Ser Tyr Asn  
 100 105 110  
 Trp Leu Gly Thr Phe Asn Pro Ala Ala Ile Leu Gly  
 115 120

<210> 280  
 <211> 380  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 280  
 Met Asp Thr Asp Asn Lys Leu Phe Asn Val Gly Val Leu Leu Val Ala  
 1 5 10 15  
 Thr Leu Val Val Ala Lys Leu Ile Ser Ala Leu Leu Ile Pro Arg Ser  
 20 25 30  
 Gly Lys Arg Leu Pro Pro Val Val Arg Thr Trp Pro Val Val Gly Gly  
 35 40 45  
 Leu Leu Arg Phe Leu Lys Gly Pro Met Val Met Leu Arg Glu Glu Tyr  
 50 55 60  
 Pro Lys Leu Gly Ser Val Phe Thr Leu Asn Leu Leu Asn Lys Lys Ile  
 65 70 75 80  
 Thr Phe Phe Ile Gly Pro Glu Val Ser Ala His Phe Phe Lys Ala Ser  
 85 90 95  
 Glu Ser Asp Leu Ser Gln Gln Glu Val Tyr Gln Phe Asn Val Pro Thr  
 100 105 110  
 Phe Gly Pro Gly Val Val Phe Asp Val Asp Tyr Thr Ile Arg Gln Glu  
 115 120 125  
 Gln Phe Arg Phe Phe Thr Glu Ala Leu Arg Ile Asn Lys Leu Lys Gly  
 130 135 140  
 Tyr Val Asn Gln Met Val Met Glu Ala Glu Asp Tyr Phe Ser Lys Trp  
 145 150 155 160  
 Gly Asp Ser Gly Glu Val Asp Leu Lys Tyr Glu Leu Glu His Leu Thr  
 165 170 175  
 Ile Leu Thr Ala Ser Arg Cys Leu Leu Gly Arg Glu Val Arg Glu Lys  
 180 185 190  
 Leu Phe Asp Asp Val Ser Ala Leu Phe His Asp Leu Asp Asn Gly Met  
 195 200 205  
 Leu Pro Ile Ser Val Ile Phe Pro Tyr Leu Pro Ile Pro Ala His His  
 210 215 220  
 Arg Arg Asp Lys Ala Arg Lys Lys Leu Ser Glu Ile Phe Ala Asn Ile  
 225 230 235 240  
 Ile Ser Ser Arg Lys Cys Ala Gly Lys Ser Glu Glu Asp Met Leu Gln  
 245 250 255  
 Cys Phe Ile Asp Ser Lys Tyr Lys Asn Gly Arg Pro Thr Thr Glu Ala  
 260 265 270  
 Glu Val Thr Gly Leu Leu Ile Ala Ala Leu Phe Ala Gly Gln His Thr  
 275 280 285  
 Ser Ser Ile Thr Ser Val Trp Thr Gly Ala Tyr Leu Leu Thr Asn Lys  
 290 295 300  
 Lys Tyr Leu Ser Ala Val Ser Asn Glu Gln Lys His Leu Met Glu Lys  
 305 310 315 320  
 His Gly Asn Asn Val Asp His Asp Val Leu Ser Glu Met Asp Val Leu  
 325 330 335  
 Tyr Arg Ser Ile Lys Glu Ala Leu Arg Leu His Pro Pro Leu Ile Met  
 340 345 350  
 Leu Leu Arg Ser Ser His Ser Asp Phe Ser Val Lys Thr Arg Asp Gly  
 355 360 365  
 Lys Glu Tyr Glu Val Gly Glu Val Ser Val Leu Pro  
 370 375 380

<210> 281  
 <211> 177  
 <212> PRT  
 <213> Eucalyptus grandis

&lt;400&gt; 281

Met Trp Lys Leu Lys Ile Gly Glu Gly Ala Asn Asp Pro Tyr Leu Phe  
 1 5 10 15  
 Ser Leu Asn Asn Phe Val Gly Arg Gln Ile Trp Glu Phe Asp Pro Glu  
 20 25 30  
 Ala Gly Thr Pro Glu Glu Arg Ala Glu Val Glu Ala Ala Arg Gln Asn  
 35 40 45  
 Phe Tyr Asn Asn Arg Phe Lys Val Arg Pro Ser Ser Asp Leu Phe Trp  
 50 55 60  
 Arg Phe Gln Phe Leu Arg Glu Lys Asn Phe Lys Gln Thr Ile Pro Pro  
 65 70 75 80  
 Val Lys Ile Glu Asp Gly Glu Asp Ile Thr Tyr Glu Lys Ala Thr Ala  
 85 90 95  
 Ala Val Lys Arg Cys Val Ser Phe Trp Ser Thr Leu Gln Ser Ser His  
 100 105 110  
 Gly His Trp Pro Ala Glu Asn Ala Gly Pro Ile Ala Phe Tyr Phe Pro  
 115 120 125  
 Pro Leu Val Met Ser Leu Tyr Val Thr Gly His Leu Asn Asn Val Phe  
 130 135 140  
 His Ala Glu His Arg Arg Glu Ile Leu Arg Tyr Ile Tyr Tyr His Gln  
 145 150 155 160  
 Asn Glu Asp Gly Gly Trp Gly Leu His Ile Glu Gly His Ser Thr Met  
 165 170 175  
 Ile

&lt;210&gt; 282

&lt;211&gt; 91

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 282

His Ala Arg Gly Leu Gly Pro Pro Pro Ile Pro Val Asp Gln Phe Ser  
 1 5 10 15  
 Leu Ala Lys Leu Val Asp Ala Ile Gln Ile Met Leu Asn Pro Gln Val  
 20 25 30  
 Lys Asn Asn Ala Asp Ala Ile Ala Lys Ala Met Glu Asn Glu Asp Gly  
 35 40 45  
 Val Ser Gly Ala Val Lys Ala Phe His Lys His Leu Pro Lys Lys Met  
 50 55 60  
 Pro Gln Pro Leu Pro Pro Thr Asp His Ser Leu Ile Asp Ser Phe  
 65 70 75 80  
 Phe Thr Gly Val Gly Lys Val Phe Gly Cys Gly  
 85 90

&lt;210&gt; 283

&lt;211&gt; 172

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 283

Trp Ile Glu Gly Arg Gly Lys Ser Val Val Cys Glu Ala Ile Ile Thr  
 1 5 10 15  
 Glu Ala Val Val Ser Lys Val Leu Lys Thr Thr Val Pro Ala Leu Leu  
 20 25 30  
 Glu Leu Asn Met Leu Lys Asn Leu Thr Gly Ser Ala Leu Ala Gly Ala  
 35 40 45  
 Met Gly Gly Phe Asn Ala His Ala Ser Asn Ile Val Ser Ala Val Phe  
 50 55 60  
 Ile Ala Thr Gly Gln Asp Pro Ala Gln Asn Ile Glu Ser Ser His Cys  
 65 70 75 80

Ile Thr Met Met Glu Ala Ser Asn Asp Gly Lys Asp Leu His Val Ser  
                             85                            90                            95  
 Val Thr Met Pro Cys Ile Glu Val Gly Thr Val Gly Gly Gly Thr Gln  
                             100                            105                            110  
 Leu Ala Ser Gln Ala Ala Cys Leu Asn Met Leu Gly Val Lys Gly Ala  
                             115                            120                            125  
 Asn Lys Glu Ser Pro Gly Ala Asn Ala Gln Thr Leu Ala Arg Ile Val  
                             130                            135                            140  
 Ala Gly Ala Val Leu Ala Gly Glu Leu Ser Leu Met Ser Ala Leu Ala  
 145                            150                            155                            160  
 Ala Gly Gln Leu Val Asn Ser His Met Lys Phe Asn  
                             165                            170

&lt;210&gt; 284

&lt;211&gt; 46

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 284

Met Ala Thr Gly Gly Gly Ala Leu Asp Leu Ala Ser Gly Met Gly Gly  
   1                            5                            10                            15  
 Asn Ile Glu Lys Glu Gln Met Leu Thr Ala Val Glu Glu Tyr Glu Lys  
                             20                            25                            30  
 Tyr His Met Tyr Tyr Gly Gly Asp Glu Gly Ser Arg Lys Ser  
                             35                            40                            45

&lt;210&gt; 285

&lt;211&gt; 137

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 285

Met Ser Lys Ala Gly Ala Met Asp Leu Ala Thr Gly Leu Gly Gly Lys  
   1                            5                            10                            15  
 Met Asp Lys Ser Asp Val Leu Ser Ala Val Asp Lys Tyr Glu Lys Tyr  
                             20                            25                            30  
 His Val Cys Tyr Gly Gly Asp Glu Glu Glu Arg Arg Ala Asn Tyr Ser  
                             35                            40                            45  
 Asp Met Val Asn Lys Tyr Tyr Asp Leu Ala Thr Ser Phe Tyr Glu Phe  
                             50                            55                            60  
 Gly Trp Gly Glu Ser Phe His Phe Ala His Arg Trp Lys Gly Glu Ser  
 65                            70                            75                            80  
 Leu Arg Glu Ser Ile Lys Arg His Glu His Phe Leu Ala Leu Gln Leu  
                             85                            90                            95  
 Gly Leu Lys Pro Gly His Lys Val Leu Asp Val Gly Cys Gly Ile Gly  
                             100                            105                            110  
 Gly Pro Leu Arg Glu Ile Ala Arg Phe Ser Ser Ala Ser Val Thr Gly  
                             115                            120                            125  
 Leu Asn Asn Asn Glu Tyr Gln Ile Thr  
                             130                            135

&lt;210&gt; 286

&lt;211&gt; 117

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 286

Phe Arg Ile Trp Phe Asp Val Pro Val Val Leu Pro Pro Leu Thr Gln  
   1                            5                            10                            15  
 Cys Phe Ala Asp Arg Ile Ser Leu Val Tyr Asp Pro His Thr Asp Glu  
                             20                            25                            30

Tyr Tyr Asn Ala Pro Gly Val Glu Thr Arg Val Pro Tyr Phe Gly Ser  
 35 40 45  
 Thr Glu Gly Met Lys Tyr Leu Asp Pro Cys Phe Lys Tyr Ile Thr Pro  
 50 55 60  
 Tyr Met Ser Ser Leu Val Lys Ser Leu Glu Asp Val Gly Tyr Val Asp  
 65 70 75 80  
 Gly Lys Ser Leu Phe Gly Ala Pro Tyr Asp Phe Arg Tyr Gly Pro Gly  
 85 90 95  
 Thr Lys Ser Ser Val Gly Ala Lys Tyr Leu Glu Asn Leu Arg Lys  
 100 105 110  
 Leu Val Glu Glu Ala  
 115

<210> 287  
 <211> 27  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 287  
 Gly Tyr Trp Asn Thr Met Asp Ile Ala His Asp Arg Ala Gly Phe Tyr  
 1 5 10 15  
 Ile Cys Trp Gly Cys Leu Val Trp Val Pro Ser  
 20 25

<210> 288  
 <211> 158  
 <212> PRT  
 <213> Pinus radiata

<400> 288  
 Phe Ala Val Val Gly Pro Leu Gln Leu Thr Ser Tyr Pro Leu Ile Lys  
 1 5 10 15  
 Leu Val Gly Ile Arg Thr Gly Leu Pro Leu Pro Ser Leu Trp Glu Ile  
 20 25 30  
 Phe Ala Gln Leu Ala Val Tyr Phe Met Val Glu Asp Tyr Gly Asn Tyr  
 35 40 45  
 Trp Ile His Arg Trp Leu His Cys Lys Trp Gly Tyr Glu Lys Ile His  
 50 55 60  
 His Val His His Glu Phe Thr Ala Pro Met Gly Phe Ala Ala Pro Tyr  
 65 70 75 80  
 Ala His Trp Ser Glu Val Leu Ile Leu Gly Ile Pro Thr Phe Val Gly  
 85 90 95  
 Pro Ala Ile Ala Pro Gly His Met Ile Thr Phe Trp Cys Trp Val Val  
 100 105 110  
 Leu Arg Gln Val Glu Ala Ile Glu Thr His Ser Gly Tyr Asp Phe Pro  
 115 120 125  
 Trp Thr Leu Thr Lys Leu Ile Pro Phe Tyr Gly Gly Ala Glu Tyr His  
 130 135 140  
 Asp Tyr His His Tyr Val Gly Gly Gln Ser Gln Ser Asn Phe  
 145 150 155

<210> 289  
 <211> 113  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 289  
 Pro Ser Leu Trp Glu Ile Leu Ala Gln Leu Leu Val Tyr Phe Leu Ile  
 1 5 10 15  
 Glu Asp Tyr Thr Asn Tyr Trp Leu His Arg Leu Leu His Cys Lys Trp  
 20 25 30

Gly Tyr Glu Lys Ile His Ser Val His His Glu Tyr Ser Ala Pro Ile  
           35                          40                          45  
 Gly Phe Ala Ala Pro Tyr Ala His Trp Ala Glu Val Leu Ile Leu Gly  
           50                          55                          60  
 Ile Pro Ser Phe Leu Gly Pro Ala Ile Val Pro Gly His Met Ile Thr  
 65                          70                          75                          80  
 Leu Trp Leu Trp Ile Ala Leu Arg Gln Ile Glu Ala Ile Asp Tyr Ser  
                           85                          90                          95  
 Gln Arg Val Arg Ile Ala Leu Glu Ser Tyr Glu Val His Ser Ile Leu  
                           100                          105                          110  
 Trp

<210> 290  
 <211> 128  
 <212> PRT  
 <213> Eucalyptus grandis

<400> 290  
 Gly Tyr Gly Ser Met Val Gln Asn Cys Val Lys Ala Arg Ser Leu Leu  
   1                          5                          10                          15  
 Ser Lys Leu Gly Ile Glu Val Thr Val Ala Asp Ala Arg Phe Cys Lys  
           20                          25                          30  
 Pro Leu Asp Ile Gly Leu Leu Arg Glu Leu Cys Glu Asn His Ala Phe  
           35                          40                          45  
 Leu Val Thr Val Glu Glu Gly Ser Ile Gly Gly Phe Gly Ser His Val  
           50                          55                          60  
 Ala Gln Phe Ile Ala Leu Asp Gly Arg Leu Asp Gly Arg Ile Lys Trp  
 65                          70                          75                          80  
 Arg Pro Ile Val Leu Pro Asp Ala Tyr Val Glu His Thr Ser Pro Asn  
                           85                          90                          95  
 Glu Gln Leu Ser Leu Ala Gly Leu Thr Gly His His Ile Ala Ala Thr  
                           100                          105                          110  
 Val Leu Ser Leu Leu Gly Arg Thr Arg Glu Ala Leu Leu Leu Met Cys  
           115                          120                          125

<210> 291  
 <211> 109  
 <212> PRT  
 <213> Pinus radiata

<400> 291  
 Met Ala Val Val Ser Ala Pro Gly Lys Val Leu Ile Thr Gly Ala  
   1                          5                          10                          15  
 Tyr Leu Ile Leu Glu Lys Pro Asn Pro Gly Leu Val Leu Thr Thr Thr  
           20                          25                          30  
 Ala Arg Phe Tyr Ala Ile Val Lys Pro Leu Arg Thr Ser Thr Asp Ser  
           35                          40                          45  
 Ser Ser Trp Ala Trp Leu Trp Thr Asp Val Lys Leu Thr Ser Pro Gln  
           50                          55                          60  
 Leu Ala Lys Glu Ala Ile Tyr Lys Leu Ser Leu Lys Thr Leu Ser Leu  
 65                          70                          75                          80  
 Gln Asn Val Ala Ser Ser Ser Ser Asn Gly Asn Pro Phe Val Glu Gln  
                           85                          90                          95  
 Ala Val Gln Phe Ala Val Ala Ala Ala Lys Glu Ala Phe  
           100                          105

<210> 292  
 <211> 107  
 <212> PRT  
 <213> Eucalyptus grandis

&lt;400&gt; 292

Met Ala Gly Glu Trp Ile Leu Thr Leu Thr Ala Gln Thr Pro Thr Asn  
 1 5 10 15  
 Ile Ala Val Ile Lys Tyr Trp Gly Lys Arg Asp Glu Ser Leu Ile Leu  
 20 25 30  
 Pro Val Asn Asp Ser Ile Ser Val Thr Leu Asp Pro Gly His Leu Cys  
 35 40 45  
 Thr Thr Thr Thr Val Ala Val Ser Pro Ala Phe Glu Gln Asp Arg Met  
 50 55 60  
 Trp Leu Asn Gly Lys Glu Ile Ser Leu Ser Gly Asp Arg Phe Gln Ser  
 65 70 75 80  
 Cys Leu Arg Glu Ile Arg Ala Arg Ala Thr Asp Val Glu Asn Lys Glu  
 85 90 95  
 Lys Gly Ile Lys Ile Ser Lys Lys Asp Trp Glu  
 100 105

&lt;210&gt; 293

&lt;211&gt; 148

&lt;212&gt; PRT

&lt;213&gt; Pinus radiata

&lt;400&gt; 293

Pro Leu Thr Leu Leu Leu Ala Asn Thr Trp Ala Ser Ser Ala Ile Val  
 1 5 10 15  
 Ser Arg Arg Val Ser Leu Phe Val Ala Cys Ser Thr Thr Val Val Ser  
 20 25 30  
 Arg Ser Phe Ser Lys Ser Cys Ser Gly Ala Ile Pro Arg Lys Pro Lys  
 35 40 45  
 Ser Ala His Pro Ala Leu Thr Gly Ser Arg Thr Cys Phe Ser Arg Asn  
 50 55 60  
 Pro Ile Val Arg Asn Leu Ile Gly Ser Ala Ser Lys Met Gly Ala Thr  
 65 70 75 80  
 Val Glu Asp Thr Thr Met Asp Ala Val Gln Arg Arg Leu Met Phe Glu  
 85 90 95  
 Asp Glu Cys Ile Leu Val Asp Glu Glu Asp His Val Ile Gly His Asp  
 100 105 110  
 Ser Lys Tyr Asn Cys His Leu Met Glu Lys Ile Glu Ser Glu Asn Leu  
 115 120 125  
 Leu His Arg Ala Phe Ser Val Phe Leu Phe Asn Thr Lys Tyr Glu Leu  
 130 135 140  
 Leu Leu Gln Gln  
 145

&lt;210&gt; 294

&lt;211&gt; 137

&lt;212&gt; PRT

&lt;213&gt; Eucalyptus grandis

&lt;400&gt; 294

Pro Leu Leu Leu Leu Leu Leu Arg Tyr Pro Ser Pro Leu Pro Pro  
 1 5 10 15  
 Arg Pro Ser Leu Ser Leu Cys Arg Ser Thr Ala Met Ala Asp Gly Ala  
 20 25 30  
 Asp Ala Gly Met Asp Ala Val Gln Arg Arg Leu Met Phe Glu Asp Glu  
 35 40 45  
 Cys Ile Leu Val Asp Glu Asn Asp Asn Val Val Gly His Glu Ser Lys  
 50 55 60  
 Tyr Asn Cys His Leu Met Glu Lys Ile Glu Ser Leu Asn Leu Leu His  
 65 70 75 80  
 Arg Ala Phe Ser Val Phe Leu Phe Asn Ser Lys Tyr Glu Leu Leu Leu



85 90 95  
 Gln Gln Arg Ser Ala Thr Lys Val Thr Phe Pro Leu Val Trp Thr Asn  
 100 105 110  
 Thr Cys Cys Ser His Pro Leu Tyr Arg Glu Ser Glu Leu Ile Ala Glu  
 115 120 125  
 Asn Ala Leu Gly Ala Arg Asn Ala Ala  
 130 135

<210> 295  
 <211> 136  
 <212> PRT  
 <213> Pinus radiata

<400> 295  
 Ala Gly Glu Asn Leu Asp Asn His Val Asp Val Lys Asn Ile Leu Val  
 1 5 10 15  
 Gln Met Gly Thr Tyr Phe Gln Val Gln Asp Asp Tyr Leu Asp Cys Phe  
 20 25 30  
 Gly Asp Pro Glu Val Ile Gly Lys Ile Gly Thr Asp Ile Glu Asp Phe  
 35 40 45  
 Lys Cys Ser Trp Leu Val Val Gln Ala Leu Glu Arg Ala Asn Glu Ser  
 50 55 60  
 Gln Leu Gln Arg Leu Tyr Ala Asn Tyr Gly Lys Thr Asp Pro Ser Cys  
 65 70 75 80  
 Val Ala Glu Val Lys Ala Val Tyr Arg Asp Leu Gly Ile Gln Asp Val  
 85 90 95  
 Phe Phe Glu Tyr Glu Arg Thr Ser Tyr Lys Glu Leu Ile Ser Ser Ile  
 100 105 110  
 Glu Ala Gln Glu Asn Glu Ser Leu Gln Leu Val Leu Lys Ser Phe Leu  
 115 120 125  
 Gly Lys Ile Tyr Lys Arg Gln Lys  
 130 135

<210> 296  
 <211> 83  
 <212> PRT  
 <213> Pinus radiata

<400> 296  
 Met Gly Glu Ser Glu Glu Ser Leu Gly Ala Gly Ser Asn Leu Lys Ser  
 1 5 10 15  
 Ala Ala Val Leu Glu Gln Ala Lys Lys His Leu Ala Thr Asp Ala Ala  
 20 25 30  
 Gln Asp Leu Lys Lys Lys Ile Gly Leu Val Tyr Gln Leu Asn Ile Ser  
 35 40 45  
 Pro Lys Lys Ile Gly Ile Ala Glu Glu Val Phe Val Val Asp Leu Lys  
 50 55 60  
 Asn Gly Lys Val Thr Lys Gly Pro Tyr Glu Gly Lys Pro Asp Ala Thr  
 65 70 75 80  
 Phe Ser Phe

<210> 297  
 <211> 156  
 <212> PRT  
 <213> Pinus radiata

<400> 297  
 Asp Thr Ser Lys Arg Arg Met Glu Glu Ile Asn Gly Asp Asn Ala Val  
 1 5 10 15  
 Arg Arg Ser Cys Phe Pro Pro Gly Phe Met Phe Gly Ile Ala Thr Ser

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      20      25      30
Ala Tyr Gln Cys Glu Gly Ala Ala Asn Glu Gly Gly Lys Gly Pro Ser
      35      40      45
Ile Trp Asp Ser Phe Ser Arg Thr Pro Gly Lys Ile Leu Asp Gly Ser
      50      55      60
Asn Gly Asp Val Ala Val Asp Gln Tyr His Arg Tyr Lys Glu Asp Val
      65      70      75      80
Lys Leu Met Lys Asp Met Gly Val Asp Thr Tyr Arg Phe Ser Leu Ser
      85      90      95
Trp Pro Arg Ile Phe Pro Lys Gly Lys Gly Glu Ile Asn Glu Glu Gly
      100      105      110
Val Ala Tyr Tyr Asn Asn Leu Ile Asn Glu Leu Leu Gln Asn Gly Ile
      115      120      125
Gln Ala Ser Val Thr Leu Phe His Trp Asp Thr Pro Gln Ser Leu Glu
      130      135      140
Asp Glu Tyr Gly Gly Phe Leu Arg Pro Thr Ile Val
      145      150      155

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<210> 298  
 <211> 115  
 <212> PRT  
 <213> Pinus radiata

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      <400> 298
Gly Val Met Ala Gly Ile Pro Val Leu Arg Pro Phe Cys Ile Cys Leu
      1      5      10      15
Leu Ser Val Tyr Met Leu His Ile Val Ala Ala Val Ala Ser Pro Arg
      20      25      30
Leu Gly Arg Ser Ser Phe Pro Arg Gly Phe Lys Phe Gly Ala Gly Ser
      35      40      45
Ser Ala Tyr Gln Ala Glu Gly Ala Ala His Glu Gly Gly Lys Gly Pro
      50      55      60
Ser Ile Trp Asp Thr Phe Ser His Thr Pro Gly Lys Ile Ala Asp Gly
      65      70      75      80
Lys Asn Gly Asp Val Ala Val Asp Gln Tyr His Arg Tyr Lys Glu Asp
      85      90      95
Val Gln Leu Leu Lys Tyr Met Gly Met Asp Val Tyr Arg Phe Ser Ile
      100      105      110
Ser Trp Ser
      115

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<210> 299  
 <211> 127  
 <212> PRT  
 <213> Pinus radiata

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      <400> 299
Gln Arg Leu Val Ser Met Ala Leu Thr Val Glu Ala Pro Ala Ala Leu
      1      5      10      15
His Leu Gln Glu Glu Ser Glu Asn Val Lys Glu Ile Ser Arg Asp
      20      25      30
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Thr Phe Ser Tyr Thr Pro Gly Lys Ile Ile Asp Gly Arg Asn Gly Asp
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## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification <sup>7</sup> :</b> <b>C12N 5/10, 9/00, 15/29, 15/63</b>	<b>A3</b>	<b>(11) International Publication Number:</b> <b>WO 00/36081</b> <b>(43) International Publication Date:</b> 22 June 2000 (22.06.00)
<b>(21) International Application Number:</b> PCT/NZ99/00219 <b>(22) International Filing Date:</b> 16 December 1999 (16.12.99)  <b>(30) Priority Data:</b> 09/215,504 17 December 1998 (17.12.98) US 60/146,441 29 July 1999 (29.07.99) US  <b>(71) Applicants (for all designated States except US):</b> GENESIS RE-SEARCH AND DEVELOPMENT CORPORATION LIMITED [NZ/NZ]; 1 Fox Street, Parnell, Auckland (NZ). FLETCHER CHALLENGE FORESTS LIMITED [NZ/NZ]; 585 Great South Road, Penrose, Auckland (NZ).  <b>(72) Inventor; and</b> <b>(75) Inventor/Applicant (for US only):</b> HAVUKKALA, Ilkka, Jaakko [FI/NZ]; 3/121 Atkin Avenue, Mission Bay, Auckland (NZ).  <b>(74) Agents:</b> BENNETT, Michael, Roy et al.; West-Walker Bennett, Mobil on the Park, 157 Lambton Quay, Wellington (NZ).		<b>(81) Designated States:</b> AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).  <b>Published</b> <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>  <b>(88) Date of publication of the international search report:</b> 3 August 2000 (03.08.00)
<b>(54) Title:</b> MATERIALS AND METHODS FOR THE MODIFICATION OF ISOPRENOID CONTENT, COMPOSITION AND METABOLISM  <b>(57) Abstract</b>  Novel isolated polynucleotides associated with plant isoprenoid biosynthetic pathways are provided, together with genetic constructs comprising such sequences. Methods for the modulation of the content, structure and metabolism of polypeptides involved in an isoprenoid biosynthetic pathway in target organisms are also disclosed, the methods comprising incorporating one or more of the polynucleotides or genetic constructs of the present invention into the genome of a target organism. Modulation of the content, structure and metabolism of such polypeptides produces modifications in the content, structure and metabolism of isoprenoids in the target organism.		

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## INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

**A. CLASSIFICATION OF SUBJECT MATTER**Int. Cl. <sup>7</sup>: C12N 5/10, 9/00, 15/29, 15/63


According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**Minimum documentation searched (classification system followed by classification symbols)  
IPC7Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched  
See Databases below.Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)  
Details in Supplemental Box V.**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GenBank accession AJ011840 submitted 7 October 1998 by Clastre M.	1-29
X	GenBank accession AF019383 submitted 14 August 1997 by Lange BM et al.	1-29
X	GenBank accession Y15782 submitted 4 December 1997 by Camara B.	1-29
X	GenBank accession Y14333 submitted 28 July 1997 by Camara B.	1-29
X	GenBank accession AB003156 submitted 2 May 1997 by Suzuki H.	1-29

☒ Further documents are listed in the continuation of Box C ☐ See patent family annex

* Special categories of cited documents:	
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"E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
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Date of the actual completion of the international search  
5 June 2000Date of mailing of the international search report  
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# INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GenBank accession D78130 submitted 12 October 1995 by Sakakibara J et al.	1-29
X	GenBank accession AF061285 submitted 24 April 1998 by Back K et al.	1-29
X	GenBank accession U87908 submitted 31 January 1997 by Bohlmann J et al.	1-29
X	GenBank accession U87909 submitted 31 January 1997 by Bohlmann J et al.	1-29
X	GenBank accession AF006193 submitted 30 May 1997 by Bohlmann J et al.	1-29
X	GenBank accession U92266 submitted 5 March 1997 by Steele CL et al.	1-29
X	GenBank accession AF006195 submitted 30 May 1997 by Bohlmann J et al.	1-29
X	GenBank accession U60542 submitted 12 June 1996 by Kollipara KP et al.	1-29
X	GenBank accession L10390 submitted 22 September 1993 by Burnett RJ et al.	1-29
X	GenBank accession X54657 submitted 29 August 1990 by Chye ML.	1-29
X	GenBank accession U72146 submitted 21 September 1996 by Maldonado-Mendoza IE and Nessler CL.	1-29
X	GenBank accession X68652 submitted 7 October 1992 by Bach TJ.	1-29
X	GenBank accession X68651 submitted 7 October 1992 by Bach TJ.	1-29
X	GenBank accession X54659 submitted 29 August 1990 by Chye ML et al.	1-29
X	GenBank accession X15032 submitted 18 April 1989 by Caelles C.	1-29
X	GenBank accession M96068 submitted 27 April 1993 by Maldonado-Mendoza IE et al.	1-29
X	GenBank accession X96429 submitted 5 March 1996 by Chen XY et al.	1-29
X	GenBank accession U27535 submitted 23 May 1995 by Chen XY et al.	1-29
X	GenBank accession AB009029 submitted 20 November 1997 by Kushiro T.	1-29
X	GenBank accession AB009031 submitted 20 November 1997 by Kushiro T.	1-29
X	GenBank accession D89619 submitted 28 November 1996 by Shibuya M.	1-29
X	GenBank accession U02555 submitted 15 October 1993 by Matsuda SP.	1-29
X	GenBank accession U74319 submitted 15 October 1996 by Bak S et al.	1-29
X	GenBank accession Y09291 submitted 6 November 1996 by Weerck-Reichhart.	1-29



# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/NZ99/00219

C (Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GenBank accession AB014057 submitted 15 May 1998 by Kushiro T et al.	1-29
X	GenBank accession AB00930 submitted 20 November 1997 by Kushiro T.	1-29
X	GenBank accession Z83833 submitted 10 January 1997 by Warnecke D.	1-29
X	GenBank accession Z83832 submitted 10 January 1997 by Warnecke D.	1-29
X	GenBank accession U81312 submitted 7 December 1996 by Benveniste P.	1-29
X	GenBank accession U81313 submitted 7 December 1996 by Benveniste P.	1-29
X	GenBank accession AF045570 submitted 30 January 1998 by Tong Y and Nes WD.	1-29
X	GenBank accession U79669 submitted 25 November 1996 by Grebenok RJ et al.	1-29
X	GenBank accession U43683 submitted 20 December 1995 by Clouse JA.	1-29
X	GenBank accession U60205 submitted 6 June 1996 by Kaplan J and Li L.	1-29
X	GenBank accession U93162 submitted 11 March 1997 by Herrmann K.	1-29
X	GenBank accession D50559 submitted 15 May 1995 by Uwebe K.	1-29
X	GenBank accession U27099 submitted 12 May 1995 by Mandel MA et al.	1-29
X	GenBank accession Y14325 submitted 24 July 1997 by Cordier H.	1-29
X	GenBank accession U53706 submitted 6 April 1996 by Jeng CJ and Schweitzer ES.	1-29
X	GenBank accession U49260 submitted 15 February 1996 by Toth MJ et al.	1-29
X	GenBank accession Y17593 submitted 17 June 1998 by Cordier H.	1-29
X	GenBank accession Y09292 submitted 6 November 1996 by Werck-Reichhart D.	1-29
X	GenBank accession U50201 submitted 28 February 1996 by Poulton JE and Jurk S.	1-29
X	GenBank accession AF072736 submitted 16 June 1998 by Dharmawardhana D et al.	1-29
X	GenBank accession X56734 submitted 19 November 1990 by Hughes MA.	1-29
X	GenBank accession D83177 submitted 19 January 1996 by Inoue K.	1-29
X	GenBank accession U39228 submitted 23 October 1995 by Wiersma PA.	1-29
X	GenBank accession U26025 submitted 2 May 1995 by Zheng L and Poulton JE.	1-29

**INTERNATIONAL SEARCH REPORT**

International application No.

**PCT/NZ99/00219**

<b>C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT</b>		
<b>Category*</b>	<b>Citation of document, with indication, where appropriate, of the relevant passages</b>	<b>Relevant to claim No.</b>
X	GenBank accession AB017026 submitted 20 August 1998 by Snider J et al.	1-29
X	GenPept accession CAA03409 submitted 21 August 1996 by Chenivresse X et al.	1-29
X	GenPept accession CAA76803 submitted 17 June 1998 by Cordier H.	1-29
X	AU, A 24637/99 (WASHINGTON STATE UNIVERSITY RESEARCH FOUNDATION) 21 January 1999 A01H 5/00, C07K 14/415, C12N 1/00, 5/04, 5/06, 9/00, 15/29, 15/52, 15/74, 15/79, 15/82, 15/84. See entire document.	1-29

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

**Box I** Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos :  
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☒ Claims Nos : 1, part (11); 1, part (12); 2; 26, part (7); 26, part (8).  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:  
These claims refer to polynucleotide or polypeptide sequences comprising 40, 20 or 10 contiguous residues of sequences provided in SEQ. ID. NOs: 1-53, 78-286, 288-304. The scope of the claims encompasses many sequence fragments and it is not economically viable to search all possible combinations.
3. ☐ Claims Nos :  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a)

**Box II** Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

Continued in Supplemental Box

1. ☒ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

**Supplemental Box I**

(To be used when the space in any of Boxes I to VIII is not sufficient)

**Continuation of Box No: II**

The International Searching Authority has found that there are 37 separate inventions, wherein a single enzyme or protein type provides the special technical feature.

1. Nucleic and amino acid sequences SEQ. ID. NOs: 1, 252 encoding acetylcholinesterase precursor, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
2. Nucleic and amino acid sequences SEQ. ID. NOs: 2, 253 encoding deoxyxylulosephosphate synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
3. Nucleic and amino acid sequences SEQ. ID. NOs: 3, 4, 44, 254, 255, 295 encoding geranyltranstransferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
4. Nucleic and amino acid sequences SEQ. ID. NOs: 5, 6, 256, 266 encoding farnesyltranstransferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
5. Nucleic and amino acid sequences SEQ. ID. NOs: 7, 154, 258, 241 encoding squalene synthetase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
6. Nucleic and amino acid sequences SEQ. ID. NOs: 8-10, 155-157, 259-261, 242-244 encoding squalene monooxygenase. DNA probes or primers therefrom, transgenic cells and constructs containing the sequences. and methods of modulating biosynthesis of isoprenoid content and metabolism.
7. Nucleic and amino acid sequences SEQ. ID. NOs: 11, 82, 83, 262, 169, 170 encoding geranylgeranyl-diphosphate geranylgeranyltransferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
8. Nucleic and amino acid sequences SEQ. ID. NOs: 12, 263 encoding trichodiene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
9. Nucleic and amino acid sequences SEQ. ID. NOs: 13, 25-27, 84-88, 95, 115-118, 264, 276-278, 171-175, 182, 202-205 encoding pinene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
10. Nucleic and amino acid sequences SEQ. ID. NOs: 14, 89, 90, 265, 176, 177 encoding abietadine synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.

Continued in Supplemental Box II

**Supplemental Box II**

(To be used when the space in any of Boxes I to VIII is not sufficient)

**Continuation of Supplemental Box I**

11. Nucleic and amino acid sequences SEQ. ID. NOs 15, 32, 91-94, 96-98, 131-135, 266, 283, 178-181, 183-185, 218-222 encoding hydroxymethylglutaryl-CoA reductase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
12. Nucleic and amino acid sequences SEQ. ID. NOs: 16-18, 99-102, 267-269, 186-189 encoding myrcene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
13. Nucleic and amino acid sequences SEQ. ID. NOs: 19, 20, 26, 27, 103, 107, 108, 277, 278, 270, 271, 190, 194, 195 encoding limonene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
14. Nucleic and amino acid sequences SEQ. ID. NOs: 21-23, 109-111, 272-274, 196-198 encoding cadinene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
15. Nucleic and amino acid sequences SEQ. ID. NOs: 24, 114, 275, 201 encoding bisabolene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
16. Nucleic and amino acid sequences SEQ. ID. NOs: 28, 119-122, 279, 206-209 encoding cycloartenol synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
17. Nucleic and amino acid sequences SEQ. ID. NOs: 29, 124-126, 280, 211-213 encoding obtusifolioside demethylase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
18. Nucleic and amino acid sequences SEQ. ID. NOs: 30, 281 encoding lupeol synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
19. Nucleic and amino acid sequences SEQ. ID. NOs: 31, 158, 159, 282, 245, 246 encoding udp-glucose:sterol glucosyltransferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
20. Nucleic and amino acid sequences SEQ. ID. NOs: 33, 34, 160-162, 284, 285, 247-249 encoding sterolmethyltransferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.

Continued in Supplemental Box III

**Supplemental Box III**

(To be used when the space in any of Boxes I to VIII is not sufficient)

**Continuation of Supplemental Box II**

21. Nucleic and amino acid sequences SEQ. ID. NOs: 35, 136, 286, 223 encoding lecithin:cholesterol acyl transferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
22. Nucleic and amino acid sequences SEQ. ID. NOs: 36, 137, 287, 224 encoding sterol delta-7 reductase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
23. Nucleic and amino acid sequences SEQ. ID. NOs: 37, 38, 138-140, 288, 289, 225-227 encoding methyl sterol oxidase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
24. Nucleic and amino acid sequences SEQ. ID. NOs: 39, 290 encoding deoxyxylulosephosphate synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
25. Nucleic and amino acid sequences SEQ. ID. NOs: 40, 291 encoding phosphomevalonate kinase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
26. Nucleic and amino acid sequences SEQ. ID. NOs: 41, 50, 141, 142, 146, 292, 301, 228, 229, 233 encoding diphosphomevalonate decarboxylase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
27. Nucleic and amino acid sequences SEQ. ID. NOs: 42, 43, 143, 293, 294, 230 encoding isopentenyl-diphosphate delta-isomerase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
28. Nucleic and amino acid sequences SEQ. ID. NOs: 45, 296 encoding estradiol dehydrogenase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
29. Nucleic and amino acid sequences SEQ. ID. NOs: 46-49, 144, 145, 297-300, 231-232 encoding furostanol glucosidase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
30. Nucleic and amino acid sequences SEQ. ID. NOs: 51, 52, 147-153, 302, 303, 234-240 encoding oxysterol-binding protein, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.

Continued in Supplemental Box IV

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

**Supplemental Box IV**

(To be used when the space in any of Boxes I to VIII is not sufficient)

**Continuation of Supplemental Box III**

31. Nucleic and amino acid sequences SEQ. ID. NOs: 53, 304 encoding sterol carrier protein. DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
32. Nucleic and amino acid sequences SEQ. ID. NOs: 78, 79, 127-130, 165, 166, 214-217 encoding sterol 14-demethylase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
33. Nucleic and amino acid sequences SEQ. ID. NOs: 82, 83, 169, 170 encoding geranylgeranyl diphosphate, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
34. Nucleic and amino acid sequences SEQ. ID. NOs: 104-106, 164, 191-193, 251 encoding CXPS/transketolase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
35. Nucleic and amino acid sequences SEQ. ID. NOs: 112, 113, 199, 200 encoding sabinene synthase. DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
36. Nucleic and amino acid sequences SEQ. ID. NOs: 123, 210 encoding beta-amyrin synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
37. Nucleic and amino acid sequences SEQ. ID. NOs: 163, 250 encoding sterol desaturase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.

The above inventions have been allocated into the following groups for searching purposes:

- A: Inventions 1 to 6.
- B: Inventions 7 to 10.
- C: Inventions 11 and 12.
- D: Inventions 13 to 15.
- E: Inventions 16 to 20.
- F: Inventions 21 to 26.
- G: Inventions 27 to 30.
- H: Inventions 31 to 37.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

**Supplemental Box V**

(To be used when the space in any of Boxes I to VIII is not sufficient)

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used):

GenBank, EMBL, PDB Nucleic Acids, SWISS-PROT, GenPept, PIR, TREMBL - SEQ. ID. NOS: 1-53, 78-286, 288-304

WPIDS: Keywords used - acetylcholinesterase precursor, deoxyxylulosephosphate synthase, dxps, geranyltranstransferase, farnesyl diphosphate synthase, farnesyltranstransferase, farnesyl diphosphate farnesyltransferase, presqualene diphosphate, squalene synthetase, squalene monooxygenase, squalene epoxidase, geranylgeranyl diphosphate geranylgeranyltransferase, prephytoene diphosphate synthase, trichodiene synthase, pinene synthase, abietadine synthase, hydroxymethylglutaryl coa reductase, myrcene synthase, limonene synthase, cadinene synthase, bisabolene synthase, cycloartenol synthase, epoxysqualene cycloarteno cyclase, obtusifolioside demethylase, lupeol synthase, udp glucose sterol glucosyl transferase, sterol glucosyltransferase, sterolmethyltransferase, lecithin cholesterol acyl transferase, phospholipid cholesterol acyltransferase, sterol delta 7 reductase, methyl sterol oxidase, deoxyxylulosephosphate synthase, dxps, diphosphomevalonate decarboxylase, phosphomevalonate kinase, isopentenyl diphosphate delta isomerase, estradioldehydrogenase, furostanol glucosidase, oxysterol binding protein, sterol carrier protein, sterol 14 demethylase, sesquiterpene cyclase, trichodiene synthase, dxps transketolase, sabinene synthase, beta amyrin synthase, sterol desaturase, pinus radiata, p radiata, pine, or pinus, eucalyptus grandis, e grandis, eucalyptus, isoprenylation, isoprenoid



## INTERNATIONAL SEARCH REPORT

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report		Patent Family Member	
AU	24637/99	WO	9937139
END OF ANNEX			